

Query Match 100.0%; Score 55; DB 2; Length 55;
Best Local Similarity 100.0%; Pred. No. 7.8e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTACATTGACCTAGTGGCCGGGCAAGCCCGGGGCGGCACTAGTCAATCGAT 55
DB 1 GTACATTGACCTAGTGGCCGGGCAAGCCCGGGGCGGCACTAGTCAATCGAT 55

RESULT 2
US-08-892-873-19
; Sequence 19, Application US/08892873
; Patent No. 6033908
; GENERAL INFORMATION:
; APPLICANT: FALLAUX et al.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
; TITLE OF INVENTION: ADENOVIRUS TO BE USED IN GENE THERAPY
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: RAE-VENTER LAW GROUP, P.C.
; STREET: 260 SHERIDAN AVENUE, PO BOX 60039
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94306
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/892,873
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/793,170
; FILING DATE: 25-MAR-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO 97/00326
; FILING DATE: 14-JUN-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: EP 95201728.3
; FILING DATE: 26-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: EP 95201611.1
; FILING DATE: 15-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: RAE-VENTER, BARBARA
; REGISTRATION NUMBER: 32,750
; REFERENCE/DOCKET NUMBER: INGR.002.000S
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650)328-4400
; TELEFAX: (650)328-3377
; TELEX: N/A
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 55 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; US-08-892-873-19

Query Match 100.0%; Score 55; DB 3; Length 55;
Best Local Similarity 100.0%; Pred. No. 7.8e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTACATTGACCTAGTGGCCGGGCAAGCCCGGGGCGGCACTAGTCAATCGAT 55
DB 1 GTACATTGACCTAGTGGCCGGGCAAGCCCGGGGCGGCACTAGTCAATCGAT 55

RESULT 3
US-09-334-765A-19
; Sequence 19, Application US/09334765A
; Patent No. 6238893
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoebein, Robert C.
; APPLICANT: Boul, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: Van der Eb, Alex J.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO BE USED
; TITLE OF INVENTION: IN GENE THERAPY
; FILE REFERENCE: 3633.20S
; CURRENT APPLICATION NUMBER: US/09/334,765A
; CURRENT FILING DATE: 1999-06-16
; PRIOR APPLICATION NUMBER: US 08/793,170
; PRIOR FILING DATE: 1997-03-25
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 19
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/clal
; US-09-334-765A-19

Query Match 100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 7.8e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTACATTGACCTAGTGGCCGGGCAAGCCCGGGGCGGCACTAGTCAATCGAT 55
DB 1 GTACATTGACCTAGTGGCCGGGCAAGCCCGGGGCGGCACTAGTCAATCGAT 55

RESULT 4
US-09-356-575E-19
; Sequence 19, Application US/09356575E
; Patent No. 6265212
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits
; APPLICANT: Hoebein, Robert
; APPLICANT: Boul, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: van der Eb, Alex
; APPLICANT: Schouten, Govert
; TITLE OF INVENTION: PACKAGING SYSTEMS
; FILE REFERENCE: 2578-39350S
; CURRENT APPLICATION NUMBER: US/09/356,575E
; CURRENT FILING DATE: 1999-07-19
; PRIOR APPLICATION NUMBER: US 08/793,170
; PRIOR FILING DATE: 1997-03-25
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Patentin Version 3.0
; SEQ ID NO 19
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Unknown

```
; FEATURE:
; OTHER INFORMATION: Derived from Adenovirus
US-09-356-575E-19
Query Match          100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 7.8e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTACATTGACCTAGTCCCGCGGCAAAAGCCCGGCGGCACTAGGTCAATCGAT 55
DB 1 GTACATTGACCTAGTCCCGCGGCAAAAGCCCGGCGGCACTAGGTCAATCGAT 55

RESULT 5
US-09-333-820-19
; Sequence 19, Application US/09333820A
; Patent No. 6306652
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoebe, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: Van der Eb, Alex J.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO BE USED IN
; FILE REFERENCE: 3833.1US
; CURRENT APPLICATION NUMBER: US/09/333,820A
; CURRENT FILING DATE: 1999-06-15
; EARLIER APPLICATION NUMBER: US 08/793,170
; EARLIER FILING DATE: 1997-03-25
; EARLIER APPLICATION NUMBER: PCT/NL96/00244
; EARLIER FILING DATE: 1996-06-14
; EARLIER APPLICATION NUMBER: EP 95201728.3
; EARLIER FILING DATE: 1995-06-26
; EARLIER APPLICATION NUMBER: EP 95201611.1
; EARLIER FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel Wordperfect 8.0
; SEQ ID NO 19
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/clal
US-09-333-820-19

Query Match          100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 7.8e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTACATTGACCTAGTCCCGCGGCAAAAGCCCGGCGGCACTAGGTCAATCGAT 55
DB 1 GTACATTGACCTAGTCCCGCGGCAAAAGCCCGGCGGCACTAGGTCAATCGAT 55

RESULT 6
US-09-358-036-41
; Sequence 41, Application US/09358036
; Patent No. 6340595
; GENERAL INFORMATION:
; APPLICANT: Vogels, Ronald
; APPLICANT: Bout, Abraham
; APPLICANT: van Es, Helmut
; APPLICANT: Schouten, Goevert
; TITLE OF INVENTION: High Throughput Screening of Gene Function Using
; TITLE OF INVENTION: Adenoviral Libraries for Functional Genomics
; FILE REFERENCE: 21834108
; CURRENT APPLICATION NUMBER: US/09/358,036
; CURRENT FILING DATE: 1999-07-21
; EARLIER APPLICATION NUMBER: US 09/097,239
```

```
; EARLIER FILING DATE: 1995-07-25
; NUMBER OF SEQ ID NOS: 69
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 41
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: o1lgonucleotide
US-09-358-036-41

Query Match          100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 7.8e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTACATTGACCTAGTCCCGCGGCAAAAGCCCGGCGGCACTAGGTCAATCGAT 55
DB 1 GTACATTGACCTAGTCCCGCGGCAAAAGCCCGGCGGCACTAGGTCAATCGAT 55

RESULT 7
US-09-097-239-41
; Sequence 41, Application US/09097239
; Patent No. 6413776
; GENERAL INFORMATION:
; APPLICANT: VOGELS, RONALD,
; APPLICANT: BOUT, ABRAHAM,
; APPLICANT: VAN ES, HELMUT H,
; APPLICANT: SCHOUTEN, GOEVERT
; TITLE OF INVENTION: HIGH THROUGHPUT SCREENING OF GENE
; TITLE OF INVENTION: FUNCTION USING ADENOVIRAL LIBRARIES FOR FUNCTIONAL
; NUMBER OF SEQUENCES: 47
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: RAE-YENTER LAW GROUP
; STREET: PO BOX 60039
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94306
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentln Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/097,239
; FILING DATE: 12-JUN-1998
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: RAE-YENTER, BARBARA
; REGISTRATION NUMBER: 32,750
; REFERENCE/DOCKET NUMBER: INGE.008.000US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650)328-4400
; TELEFAX: (650)328-4477
; INFORMATION FOR SEQ ID NO: 41:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 55 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
US-09-097-239-41

Query Match          100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 7.8e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTACATTGACCTAGTCCCGCGGCAAAAGCCCGGCGGCACTAGGTCAATCGAT 55
DB 1 GTACATTGACCTAGTCCCGCGGCAAAAGCCCGGCGGCACTAGGTCAATCGAT 55
```

RESULT 8
US-08-793-170-20/c
Sequence 20, Application US/08793170
Patent No. 5994128
GENERAL INFORMATION:
APPLICANT: FALLAUX et al.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
ADENOVIRUS TO BE USED IN GENE THERAPY
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: RAE-VENTER LAW GROUP, P.C.
STREET: 260 SHERIDAN AVENUE, PO BOX 60039
CITY: PALO ALTO
STATE: CALIFORNIA
COUNTRY: USA
ZIP: 94306
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/793,170
FILING DATE: 25-MAR-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO 97/00326
FILING DATE: 14-JUN-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: EP 95201728.3
FILING DATE: 26-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: EP 95201611.1
FILING DATE: 15-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: RAE-VENTER, BARBARA
REGISTRATION NUMBER: 32,750
REFERENCE/DOCKET NUMBER: INGE.002.000S
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650)328-4400
TELEFAX: (650)328-3377
TELEX: N/A
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 55 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
US-08-793-170-20

Query Match 92.7%; Score 51; DB 2; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.9e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 ATTGACCTAGTCCGCCGCGCAAGCCCGGCGGCACTAGTCAATCGAT 55
DB 55 ATTGACCTAGTCCGCCGCGCAAGCCCGGCGGCGGCACTAGTCAATCGAT 5

RESULT 9
US-08-892-873-20/c
Sequence 20, Application US/08892873
Patent No. 6033908
GENERAL INFORMATION:
APPLICANT: FALLAUX et al.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
ADENOVIRUS TO BE USED IN GENE THERAPY
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: RAE-VENTER LAW GROUP, P.C.
STREET: 260 SHERIDAN AVENUE, PO BOX 60039

CITY: PALO ALTO
STATE: CALIFORNIA
COUNTRY: USA
ZIP: 94306
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/892,873
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/793,170
FILING DATE: 25-MAR-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO 97/00326
FILING DATE: 14-JUN-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: EP 95201728.3
FILING DATE: 26-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: EP 95201611.1
FILING DATE: 15-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: RAE-VENTER, BARBARA
REGISTRATION NUMBER: 32,750
REFERENCE/DOCKET NUMBER: INGE.002.000S
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650)328-4400
TELEFAX: (650)328-3377
TELEX: N/A
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 55 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
US-08-892-873-20

Query Match 92.7%; Score 51; DB 3; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.9e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 ATTGACCTAGTCCGCCGCGCAAGCCCGGCGGCACTAGTCAATCGAT 55
DB 55 ATTGACCTAGTCCGCCGCGCAAGCCCGGCGGCGGCACTAGTCAATCGAT 5

RESULT 10
US-09-334-765A-20/c
Sequence 20, Application US/09334765A
Patent No. 6238893
GENERAL INFORMATION:
APPLICANT: Fallaux, Frits J.
APPLICANT: Hoebein, Robert C.
APPLICANT: Bout, Abraham
APPLICANT: Valetio, Domenico
APPLICANT: Van der Eb, Alex J.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
ADENOVIRUS TO BE USED
FILE REFERENCE: 3833.20S
CURRENT APPLICATION NUMBER: US/09/334,765A
PRIOR FILING DATE: 1999-06-16
PRIOR APPLICATION NUMBER: US 08/793,170
PRIOR FILING DATE: 1997-03-25
PRIOR APPLICATION NUMBER: PCT/NL96/00244
PRIOR FILING DATE: 1996-06-14
PRIOR APPLICATION NUMBER: EP 95201728.3
PRIOR FILING DATE: 1995-06-26
PRIOR APPLICATION NUMBER: EP 95201611.1

```

; PRIOR FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 20
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: primer HP/claz
US-09-334-765A-20

Query Match
Best Local Similarity 92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.9e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 5 ATTGACCTAGTCCGCCGCGGCAAAAGCCGCGGCACTAGTCAATCGAT 55
Db 55 ATTGACCTAGTCCGCCGCGGCAAAAGCCGCGGCACTAGTCAATCGAT 5

RESULT 11
US-09-356-575E-20/C
; Sequence 20, Application US/09356575E
; Patent No. 6265212
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits
; APPLICANT: Hoeber, Robert
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: van der Eb, Alex
; TITLE OF INVENTION: PACKAGING SYSTEMS
; FILE REFERENCE: 2578-3335US
; CURRENT APPLICATION NUMBER: US/09/356,575E
; CURRENT FILING DATE: 1999-07-19
; PRIOR APPLICATION NUMBER: US 08/793,170
; PRIOR FILING DATE: 1997-03-25
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 20
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; OTHER INFORMATION: Derived from Adenovirus
US-09-356-575E-20

Query Match
Best Local Similarity 92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.9e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 5 ATTGACCTAGTCCGCCGCGGCAAAAGCCGCGGCACTAGTCAATCGAT 55
Db 55 ATTGACCTAGTCCGCCGCGGCAAAAGCCGCGGCACTAGTCAATCGAT 5

RESULT 12
US-09-333-820-20/C
; Sequence 20, Application US/09333820A
; Patent No. 6306552
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoeber, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
```

```

; APPLICANT: Van der Eb, Alex J.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO BE USED
; FILE REFERENCE: 3833.1US
; CURRENT APPLICATION NUMBER: US/09/333,820A
; CURRENT FILING DATE: 1999-06-15
; PRIOR APPLICATION NUMBER: US 08/793,170
; PRIOR FILING DATE: 1997-03-25
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 20
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: primer HP/claz2
US-09-333-820-20
```

```

Query Match
Best Local Similarity 92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.9e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```

Qy 5 ATTGACCTAGTCCGCCGCGGCAAAAGCCGCGGCACTAGTCAATCGAT 55
Db 55 ATTGACCTAGTCCGCCGCGGCAAAAGCCGCGGCACTAGTCAATCGAT 5
```

```

RESULT 13
US-09-358-036-42/C
; Sequence 42, Application US/09358036
; Patent No. 6340595
; GENERAL INFORMATION:
; APPLICANT: Vogels, Ronald
; APPLICANT: Bout, Abraham
; APPLICANT: van Es, Helmut
; APPLICANT: Schouten, Goevert
; TITLE OF INVENTION: High Throughput Screening of Gene Function Using
; TITLE OF INVENTION: Adenoviral Libraries for Functional Genomics
; FILE REFERENCE: 21834108
; CURRENT APPLICATION NUMBER: US/09/358,036
; CURRENT FILING DATE: 1999-07-21
; PRIOR APPLICATION NUMBER: US 09/097,239
; PRIOR FILING DATE: 1995-07-25
; NUMBER OF SEQ ID NOS: 69
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 42
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: oligonucleotide
US-09-358-036-42
```

```

Query Match
Best Local Similarity 92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.9e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```

Qy 5 ATTGACCTAGTCCGCCGCGGCAAAAGCCGCGGCACTAGTCAATCGAT 55
Db 55 ATTGACCTAGTCCGCCGCGGCAAAAGCCGCGGCACTAGTCAATCGAT 5
```

```

RESULT 14
US-09-097-239-42/C
```

```
; Sequence 42, Application US/09097239
; Patent No. 641376
; GENERAL INFORMATION:
; APPLICANT: VOGELS, RONALD,
; APPLICANT: BOOT, ABRAHAM,
; APPLICANT: VAN ES, HELMUTH HG,
; APPLICANT: SCHOUTEN, GOVERT
; TITLE OF INVENTION: HIGH THROUGHPUT SCREENING OF GENE
; TITLE OF INVENTION: FUNCTION USING ADENOVIIRAL LIBRARIES FOR FUNCTIONAL
; TITLE OF INVENTION: GENOMICS APPLICATIONS
; NUMBER OF SEQUENCES: 47
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: RAE-VENTER LAW GROUP
; STREET: PO BOX 60039
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94306
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/097,239
; FILING DATE: 12-JUN-1998
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: RAE-VENTER, BARBARA
; REGISTRATION NUMBER: 32,750
; REFERENCE/DOCKET NUMBER: INGE.008.000S
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650)328-4400
; TELEFAX: (650)328-4477
; INFORMATION FOR SEQ ID NO: 42:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 55 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
;
US-09-097-239-42
;
Query Match          92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.9e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 5 ATTGACTAGTGGCCCGGCAAGCCGGGCGGCACTAGTCATCATCGAT 55
DB 55 ATTGACTAGTGGCCCGGCAAGCCGGGCGGCACTAGTCATCATCGAT 5
```

```
RESULT 15
US-08-793-170-17
; Sequence 17, Application US/08793170
; Patent No. 5994128
; GENERAL INFORMATION:
; APPLICANT: FALAUX et al.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
; TITLE OF INVENTION: ADENOVIRUS TO BE USED IN GENE THERAPY
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: RAE-VENTER LAW GROUP, P.C.
; STREET: 260 SHERIDAN AVENUE, PO BOX 60039
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94306
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
```

```
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/793,170
; FILING DATE: 25-MAR-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO 97/00326
; FILING DATE: 14-JUN-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: EP 95201728.3
; FILING DATE: 26-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: EP 95201611.1
; FILING DATE: 15-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: RAE-VENTER, BARBARA
; REGISTRATION NUMBER: 32,750
; REFERENCE/DOCKET NUMBER: INGE.002.000S
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650)328-4400
; TELEFAX: (650)328-3377
; TELEX: N/A
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
;
US-08-793-170-17
```

```
QY 1 GTGACTTGACCTAGTGGCCCGGCAAGCCGGGCGGCACTAGTCA 49
DB 1 GTGACTTGACCTAGTGGCCCGGCAAGCCGGGCGGCACTAGTCA 49
```

```
Search completed: December 27, 2002, 06:16:59
Job time : 35 secs
```

GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 27, 2002, 00:33:34 ; Search time 170.5 Seconds
(without alignments)
726.451 Million cell updates/sec

Title: US-09-918-029-19

Perfect score: 55
1 glactaltgacctagtagtgcgc.....gcgcactagtagtcaatcagat 55

Sequence: 1 glactaltgacctagtagtgcgc.....gcgcactagtagtcaatcagat 55

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

N_Geneseq_101002:.*
1: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1980.DAT.*
2: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1981.DAT.*
3: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1982.DAT.*
4: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1983.DAT.*
5: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1984.DAT.*
6: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1985.DAT.*
7: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1986.DAT.*
8: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1987.DAT.*
9: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1988.DAT.*
10: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1989.DAT.*
11: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1990.DAT.*
12: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1991.DAT.*
13: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1992.DAT.*
14: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1993.DAT.*
15: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1994.DAT.*
16: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1995.DAT.*
17: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1996.DAT.*
18: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1997.DAT.*
19: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1998.DAT.*
20: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1999.DAT.*
21: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2000.DAT.*
22: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001A.DAT.*
23: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001B.DAT.*
24: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	55	100.0	55	18	AA748646
2	55	100.0	55	18	AA559116
3	55	100.0	55	21	AA237959
4	55	100.0	55	22	AA30232
5	55	100.0	55	24	ABK47032
6	51	92.7	55	18	AA748647
7	51	92.7	55	21	AA559117
8	51	92.7	55	24	ABK47033
9	47.4	86.2	50	18	AA748644

10	47.4	86.2	50	22	AA730231	Oligonucleotide fo
11	47.4	86.2	50	24	ABK47030	Adenovirus vectro
12	43.4	78.9	45	21	AA559131	Haipin structure-
13	43.4	78.9	45	21	AA237961	DNA molecule conta
14	43.4	78.9	45	22	AA730234	Oligonucleotide fo
15	43.4	78.9	45	24	ABK47038	Adenovirus vectro
16	43.4	78.9	50	18	AA748645	Synthetic haipin o
17	43.4	78.9	50	21	AA237958	Adenovirus vectro
18	43.4	78.9	50	24	ABK47031	Adenovirus vectro
19	42.2	76.7	55	18	AA748647	Synthetic haipin o
20	42.2	76.7	55	18	AA748646	Synthetic haipin o
21	42.2	76.7	55	21	AA559116	Oligonucleotide HP
22	42.2	76.7	55	21	AA559117	Oligonucleotide HP
23	42.2	76.7	55	21	AA237959	Adenoviral constru
24	42.2	76.7	55	22	AA730232	Oligonucleotide fo
25	42.2	76.7	55	24	ABK47032	Adenovirus vectro
26	42.2	76.7	55	24	ABK47033	Adenovirus vectro
27	39	70.9	54	21	AA237960	Adenoviral constru
28	38.6	70.2	50	18	AA748644	Synthetic haipin o
29	38.6	70.2	50	18	AA748645	Synthetic haipin o
30	38.6	70.2	50	21	AA237958	Adenoviral constru
31	38.6	70.2	50	22	AA730231	Oligonucleotide fo
32	38.6	70.2	50	24	ABK47030	Adenovirus vectro
33	38.6	70.2	50	24	ABK47031	Adenovirus vectro
34	35.4	64.4	49	21	AA237957	Adenoviral constru
35	34.6	62.9	45	21	AA259131	Haipin structure-
36	34.6	62.9	45	21	AA237961	DNA molecule conta
37	34.6	62.9	45	22	AA730234	Oligonucleotide fo
38	34.6	62.9	45	24	ABK47038	Adenovirus vectro
39	30.2	54.9	54	21	AA237960	Adenoviral constru
40	26.6	48.4	49	21	AA237957	Adenoviral constru
41	24.8	45.1	157	21	AA211855	Human secreted pro
42	23.8	43.3	8384	22	AAK82907	Human immune/hema
43	23.6	42.9	63	24	AB558790	AAV 5, ITR from vec
44	23.6	42.9	145	14	AAQ41448	AAV2 inverted term
45	23.6	42.9	145	16	AA703385	Strict inverted te

ALIGNMENTS

RESULT 1	AA748646	AA748646 standard; DNA; 55 BP.
ID	AA748646	
XX	AA748646:	
AC		
XX		
DT	21-MAY-1997 (first entry)	
XX		
DE	Synthetic haipin oligonucleotide HP/clal.	
XX		
KW	Gene therapy; vaccine; vector; adenovirus; packaging system;	
KW	haipin; pICL; ss.	
OS	Synthetic.	
XX		
PN	WO9700326-A1.	
XX		
PD	03-JAN-1997.	
XX		
PF	14-JUN-1996; 96MO-NL00244.	
XX		
PR	26-JUN-1995; 95EP-0201728.	
XX		
PR	15-JUN-1995; 95EP-0201611.	
XX		
PA	(INTER-) INTROGENE BV.	
XX	(UYLE-) RIKSGUNIV LEIDEN.	
PI	Bout A, Pallaux FJ, Hoeven RC, Valerio D, Van Der Ebba;	
XX		
DR	WPI, 1997-077531/07.	
XX		
PT	New packaging cells and nucleic acids for recombinant adenovirus -	

PT have no overlapping sequences, prevents homologous recombination;
 PT for use in gene therapy and vaccination
 XX
 PS Disclosure; Page 55; 88pp; English.
 XX
 CC Synthetic oligonucleotides HP/clal (AA748646) and HP/claz (AA748647)
 CC were used to generate a synthetic hairpin. They contain a Clal
 CC recognition site to be used for hairpin formation. The
 CC oligonucleotides were annealed and ligated into plasmid pCMV-TK,
 CC at the adenovirus inverted terminal repeat, generating
 CC PAD-CMV-hetK. This plasmid was co-transfected with Clal-digested
 CC wild-type adenovirus 4 into 911 cells. A recombinant adenovirus
 CC in which the CMV-hetK expression cassette replaced the E1 sequences
 CC was isolated.
 XX
 SQ Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;
 XX
 Query Match 100.0%; Score 55; DB 18; Length 55;
 Best Local Similarity 100.0%; Pred. No. 1.3e-10;
 Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GTACATTGACCTAGTACCGCCGCGCAAGCCGCGGCGACTAGTCAATCGAT 55
 1 GTACATTGACCTAGTACCGCCGCGCAAGCCGCGGCGACTAGTCAATCGAT 55
 DB
 RESULT 2
 AA259116
 ID AA259116 standard; DNA; 55 BP.
 XX
 AC AA259116;
 XX
 DT 11-APR-2000 (first entry)
 XX
 DE Oligonucleotide HP/clal for generating hairpin structure.
 XX
 KW Expressible nucleic acid library; gene expression; gene function;
 KW capillary formation; cell proliferation; hairpin structure; ss.
 XX
 OS Synthetic.
 XX
 PN MO964582-A2.
 XX
 PD 16-DEC-1999.
 XX
 PF 11-JUN-1999; 99WO-NL00367.
 XX
 PR 12-JUN-1998; 98US-0097239.
 XX
 PA (INTR-) INTROGENE BV.
 XX
 PI Schouten G, Vogels R, Bout A, Van Es H;
 XX
 DR WPI: 2000-097536/08.
 XX
 PT New library of expressible nucleic acids, useful for high-throughput
 PT screening of gene function, especially for identifying therapeutic
 PT molecules -
 XX
 PS Example 3; Page 156; 223pp; English.
 XX
 CC The invention relates to a library of expressible nucleic acids (NA)
 CC which contains many compartments, each comprising at least one vehicle
 CC comprising at least one NA, the vehicle being capable of efficiently
 CC introducing a NA into a cell for expression. The library is useful for
 CC determining the function of one or more nucleic acids within the
 CC library, or to screen for an expressible nucleic acid with a particular
 CC desired function. It is especially useful for high throughput screening
 CC of gene function for functional genomics applications and for screening
 CC for nucleic acids with potential therapeutic value. Cell types
 CC appropriate for selection of a particular phenotype may be useful for
 CC capillary formation and cell proliferation. Oligonucleotides
 CC AA259116-259117 were used to generate a hairpin structure in plasmid

CC PAD-CMV-hetK. The hairpin structure was used to determine if it could be
 CC used to prime reverse strand synthesis on the displaced strand after
 CC replication initiation in the adenoviral inverted terminal repeat (ITR).
 CC
 SQ Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;
 XX
 Query Match 100.0%; Score 55; DB 21; Length 55;
 Best Local Similarity 100.0%; Pred. No. 1.3e-10;
 Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GTACATTGACCTAGTACCGCCGCGCAAGCCGCGGCGACTAGTCAATCGAT 55
 1 GTACATTGACCTAGTACCGCCGCGCAAGCCGCGGCGACTAGTCAATCGAT 55
 DB
 RESULT 3
 AA237959
 ID AA237959 standard; DNA; 55 BP.
 XX
 AC AA237959;
 XX
 DT 07-FEB-2000 (first entry)
 XX
 DE Adenoviral construct generating primer Hp/clal.
 XX
 KW Adenoviral vector; replication-defective; adenovirus; ITR; hepatitis;
 KW inverted terminal repeat; encapsulation signal; gene therapy; tumor;
 KW inherited disease; cystic fibrosis; Duchenne molecular dystrophy;
 KW hypercholesterolemia; blood clotting disorder; hemophilia; restenosis;
 KW autoimmune disease; rheumatoid arthritis; PCR primer; ss.
 XX
 OS Synthetic.
 XX
 PN MO955132-A2.
 XX
 PD 04-NOV-1999.
 XX
 PF 23-APR-1999; 99WO-NL00235.
 XX
 PR 24-APR-1998; 98US-0065752.
 XX
 PA (INTR-) INTROGENE BV.
 XX
 PI Vogels R, Bout A;
 XX
 DR WPI: 2000-023229/02.
 XX
 PT New recombinant adenovirus vectors, used particularly for gene therapy
 PT for treating inherited or acquired diseases -
 XX
 PS Disclosure; Page 118; 161pp; English.
 XX
 CC The invention provides methods of producing recombinant adenoviral
 CC vectors (Adv's) for generating replication-defective adenoviruses.
 CC Generating an Adv comprises fusing 2 partially overlapping sequences
 CC nucleic acid molecules that are capable of combining with each other to
 CC allow the generation of a physically linked nucleic acid comprising at
 CC least 2 functional adenoviral inverted terminal repeats (ITRs), a
 CC functional encapsulation signal and a nucleic acid of interest. The
 CC products can be used for gene therapy for treating inherited diseases
 CC e.g. cystic fibrosis, Duchenne molecular dystrophy,
 CC hypercholesterolemia, blood clotting disorders (hemophilia) or acquired
 CC diseases such as tumors, hepatitis, (auto)immune diseases, restenosis, or
 CC rheumatoid arthritis. Sequences AA237954-960 represent primers used for
 CC PCR amplification of DNA fragments used for generation of adenoviral
 CC constructs of the invention.
 XX
 SQ Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;
 XX
 Query Match 100.0%; Score 55; DB 21; Length 55;
 Best Local Similarity 100.0%; Pred. No. 1.3e-10;
 Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTACATTGACCTAGTCCGCCCGGCAAAAGCCCGGCGGACACTAGTCAATGCAT 55
 |||||||
 DB 1 GTACATTGACCTAGTCCGCCCGGCAAAAGCCCGGCGGACACTAGTCAATGCAT 55
 |||||||

RESULT 4
 AAF30232
 ID AAF30232 standard; DNA; 55 BP.

XX AAF30232;

XX 30-APR-2001 (first entry)

XX Oligonucleotide forming hairpin structure.

XX Adenovirus; vector; gene therapy; packaging cell; hairpin; ds.

XX Synthetic.

XX Key location/Qualifiers

XX misc_feature 1..4

XX /tag- 2
 /note- "single-stranded 5' overhang"

XX misc_feature 55

XX /tag- b
 /note- "single-stranded overhang on complementary strand of sequence 5'-GTAC-3'

XX NO200105945-A2.

XX 25-JAN-2001.

XX 19-JUL-2000; 2000WO-EP07074.

XX 19-JUL-1999; 99US-0356575.

XX (INTNR-) INTROGENE BV.

XX Hoebein RC, Bout A, Valerio D, Van Der Eb AJ, Schouten G;

XX Fallaux EJ;

XX WPI; 2001-147334/15.

XX Producing recombinant adenovirus for use in gene therapy, comprises
 PT culturing cells containing adenoviral nucleic acid having an
 PT encapsidating signal and inverted terminal repeat, and lacking
 PT overlapping sequences -

XX Example; Page 39; 97pp; English.

XX The present sequence is that of an oligonucleotide formed from 2
 CC partially complementary oligonucleotides creating a hairpin
 CC structure. The oligonucleotide forms an *clat* recognition site
 CC when inserted into the *clat* site of plasmid pUC19 (see AAF30233).
 CC the hairpin could be used as a primer for reverse strand synthesis
 CC on the displaced strand after replication had started from the
 CC inverted terminal repeat (ITR) of the vector. In adenovirus
 CC infected cells, linear DNA fragments have on one terminus an
 CC adenovirus-derived ITR and at the other terminus a sequence that
 CC can anneal to the same strand, when present in single-stranded
 CC form, thereby generating a hairpin structure, and will be
 CC converted to structures with ITRs at both ends. The resulting DNA
 CC molecules will replicate by the same mechanism as the wild-type
 CC adenovirus genomes. The invention provides adenovirus vectors and
 CC packaging cell lines useful in the safe generation of *E1*-deleted
 CC recombinant adenovirus vectors for gene therapy applications.
 CC Packaging cells contain adenovirus nucleic acids having an
 CC encapsidating signal and ITR, but lack sequences that overlap with
 CC the vector, thereby preventing homologous recombination.

XX Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;

Query Match 100.0%; Score 55; DB 22; Length 55;
 Best Local Similarity 100.0%; Pred. No. 1.3e-10;
 Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTACATTGACCTAGTCCGCCCGGCAAAAGCCCGGCGGACACTAGTCAATGCAT 55
 |||||||
 DB 1 GTACATTGACCTAGTCCGCCCGGCAAAAGCCCGGCGGACACTAGTCAATGCAT 55
 |||||||

RESULT 5

XX ABR47032
 ID ABR47032 standard; DNA; 55 BP.

XX ABR47032;

XX 05-JUN-2002 (first entry)

XX Adenovirus vector pUC19ac/haw hairpin linker sequence *Hp/clat*1.

XX Adenovirus vector library; ss; linker; high throughput screening;

XX RCA; replication competent adenovirus.

XX Synthetic.

XX US6340595-B1.

XX 21-JUL-1999; 99US-0358036.

XX 12-JUN-1998; 98US-0097239.

XX (GALA-) GALAPAGOS GENOMICS NV.

XX Vogels R, Bout A, Van Es H, Schouten G;

XX WPI; 2002-224926/28.

XX Library of expressible nucleic acids, useful for determining nucleic
 PT acid function, comprises one or more adenoviral vectors capable of
 PT transfecting a host cell with the nucleic acid -

XX Example 3; Column 85; 11pp; English.

XX The invention relates to a library (I) of a multitude of unique
 CC expressible nucleic acids (NA), comprises a number of compartments
 CC (II), each consisting of one or more adenoviral vectors (III)
 CC comprising at least one unique NA of (I) in an aqueous medium, where
 CC (III) is capable of introducing the NA into a host cell (IV), is
 CC capable of expressing the product of the NA in (IV), and is deleted in
 CC a portion of the adenoviral genome necessary for replication. Also
 CC included is a method for producing the library. The library is useful for
 CC determining the function of at least one nucleic acid that is present.
 CC The library uses high throughput generation of recombinant adenoviral
 CC vector libraries containing one or more sample nucleic acids, followed by
 CC high throughput screening of the adenoviral vector libraries in a host to
 CC alter the phenotype of the host as a means of assigning a function to
 CC expression product(s) of the sample nucleic acids. The entire process
 CC lends itself to automation especially when implemented in a 96-well or
 CC other multi-well format. The high throughput screening, using a number of
 CC different *in vitro* assays, provides a means of efficiently obtaining
 CC functional information in a relatively short period of time. The
 CC member(s) of the recombinant adenoviral libraries that exhibit or induce
 CC a desired phenotype in a host *in vitro* or *in situ* are identified to
 CC reduce the libraries to a manageable number of recombinant adenoviral
 CC vectors or clones which can be tested *in vitro* in an animal model.
 CC Furthermore, the methods produce RCA-free adenoviral libraries. RCA
 CC (replication competent adenovirus) contamination throughout the libraries
 CC could become a major obstacle, especially if libraries are continuously
 CC amplified for use in multiple screening programs. Additionally, a further
 CC advantage is minimisation of the number of steps involved in the process.
 CC There is no requirement for cloning each individual adenovirus before use
 CC in a high throughput screening program. Other systems include one or more

CC steps in E. coli to achieve homologous recombination for the various
 CC adenoviral plasmids necessary for vector generation. The present
 CC sequence is a linker sequence used in the construction of the adenoviral
 CC vector library of the invention.

SO Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;

Query Match 100.0%; Score 55; DB 24; Length 55;
 Best Local Similarity 100.0%; Pred. No. 1.3e-10;
 Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTACATTCAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55
 1 GTACATTCAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55
 DB 1 GTACATTCAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55

RESULT 6
 AAT48647/C
 ID AAT48647 standard; DNA: 55 BP.

AC AAT48647;

DT 21-MAR-1997 (first entry)

DE Synthetic hairpin oligonucleotide HP/c1a2.

KW Gene therapy; vaccine; vector; adenovirus; packaging system;
 KW hairpin; PICL; ss.

OS Synthetic.

PN WO9700326-A1.

PD 03-JAN-1997.

PF 14-JUN-1996; 96WO-NL00244.

PR 26-JUN-1995; 95FP-0201728.

PR 15-JUN-1995; 95EP-0201611.

PA (INTR-) INTROGENE BV.

PA (UYLE-) RIJCKSUNIV LEIDEN.

PI Bout A, Fallaux FJ, Hoeven RC, Valerio D, Van Der EBBADJ;

DR WPI: 1997-077531/07.

PT New packaging cells and nucleic acids for recombinant adenovirus -
 PT have no overlapping sequences, prevents homologous recombination;
 PT for use in gene therapy and vaccination

PS Disclosure: Page 55; 88pp; English.

CC Synthetic oligonucleotides HP/c1a1 (AAT48646) and HP/c1a2 (AAT48647)
 CC were used to generate a synthetic hairpin. They contain a ClaI
 CC recognition site to be used for hairpin formation. The
 CC oligonucleotides were annealed and ligated into plasmid pCMV.TK,
 CC at the adenovirus inverted terminal repeat, generating
 CC pad-CMV-hcTK. This plasmid was co-transfected with ClaI-digested
 CC wild-type adenovirus 4 into 911 cells. A recombinant adenovirus
 CC in which the CMV-hcTK expression cassette replaced the EI sequences
 CC was isolated.

SO Sequence 55 BP; 9 A; 17 C; 17 G; 12 T; 0 other;

Query Match 92.7%; Score 51; DB 18; Length 55;
 Best Local Similarity 100.0%; Pred. No. 3.6e-09;
 Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 ATTGACCTAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55
 5 ATTGACCTAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55
 DB 55 ATTGACCTAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55
 XX

RESULT 7
 AAZ59117/C
 ID AAZ59117 standard; DNA: 55 BP.

AC AAZ59117;

DT 11-APR-2000 (first entry)

DE Oligonucleotide HP/c1a2 for generating hairpin structure.

KW Expressible nucleic acid library; gene expression; gene function;
 KW capillary formation; cell proliferation; hairpin structure; ss.

OS Synthetic.

PN WO964582-A2.

PD 16-DEC-1999.

PF 11-JUN-1999; 99WO-NL00367.

PR 12-JUN-1998; 98US-0097239.

PA (INTR-) INTROGENE BV.

PI Schouten G, Vogels R, Bout A, Van Es H;

DR WPI: 2000-097536/08.

PT New library of expressible nucleic acids, useful for high-throughput
 PT screening of gene function, especially for identifying therapeutic
 PT molecules

PS Example 3; Page 156; 223pp; English.

CC The invention relates to a library of expressible nucleic acids (NA)
 CC which contains many compartments, each comprising at least one vehicle
 CC comprising at least one NA, the vehicle being capable of efficiently
 CC introducing a NA into a cell for expression. The library is useful for
 CC determining the function of one or more nucleic acids within the
 CC library, or to screen for an expressible nucleic acid with a particular
 CC desired function. It is especially useful for high throughput screening
 CC of gene function for functional genomics applications and for screening
 CC for nucleic acids with potential therapeutic value. Cell types
 CC appropriate for selection of a particular phenotype may be useful for
 CC capillary formation and cell proliferation. Oligonucleotides
 CC AAZ59116-259117 were used to generate a hairpin structure in plasmid
 CC pad-CMV-hcTK. The hairpin structure was used to determine if it could be
 CC used to prime reverse strand synthesis on the displaced strand after
 CC replication initiation in the adenoviral inverted terminal repeat (ITR).

SO Sequence 55 BP; 9 A; 17 C; 17 G; 12 T; 0 other;

Query Match 92.7%; Score 51; DB 21; Length 55;
 Best Local Similarity 100.0%; Pred. No. 3.6e-09;
 Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 ATTGACCTAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55
 5 ATTGACCTAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55
 DB 55 ATTGACCTAGTCGCGCGGCAAGCCCGGGCGGCACTAGTCAATGCAT 55
 XX

RESULT 8

ABK47033/C
 ID ABK47033 standard; DNA: 55 BP.

AC ABK47033;

DT 05-JUN-2002 (first entry)

DE Adenovirus vector PIC1b/haw hairpin linker sequence HP/c1a1#2.

KW Adenovirus vector library; ss: linker; high throughput screening;
 KM RCA: replication competent adenovirus.
 XX Synthetic.
 OS
 PN US6340595-B1.
 XX
 PD 22-JAN-2002.
 XX
 PF 21-JUL-1999; 99US-0358036.
 XX
 PR 12-JUN-1998; 98US-0097239.
 XX
 PA (GALA-) GALAPAGOS GENOMICS NV.
 XX
 PI Vogels R, Bout A, Van Es H, Schouten G;
 XX
 DR WPI; 2002-224926/28.
 XX
 PT Library of expressible nucleic acids, useful for determining nucleic
 PT acid function, comprises one or more adenoviral vectors capable of
 PT transfecting a host cell with the nucleic acid -
 XX
 PS Example 3; Column 85; 11pp; English.
 XX
 CC The invention relates to a library (I) of a multitude of unique
 CC expressible nucleic acids (NA), comprises a number of compartments
 CC (II), each consisting of one or more adenoviral vectors (III)
 CC comprising at least one unique NA of (I) in an aqueous medium, where
 CC (III) is capable of introducing the NA into a host cell (IV), is
 CC capable of expressing the product of the NA in (IV), and is deleted in
 CC a portion of the adenoviral genome necessary for replication. Also
 CC included is a method for producing the library. The library is useful for
 CC determining the function of at least one nucleic acid that is present.
 CC The library uses high throughput generation of recombinant adenoviral
 CC vector libraries containing one or more sample nucleic acids, followed by
 CC high throughput screening of the adenoviral vector libraries in a host to
 CC alter the phenotype of the host as a means of assigning a function to
 CC expression product(s) of the sample nucleic acids. The entire process
 CC lends itself to automation especially when implemented in a 96-well or
 CC other multi-well format. The high throughput screening, using a number of
 CC different in vitro assays, provides a means of efficiently obtaining
 CC functional information in a relatively short period of time. The
 CC member(s) of the recombinant adenoviral libraries that exhibit or induce
 CC a desired phenotype in a host in vitro or in situ are identified to
 CC reduce the libraries to a manageable number of recombinant adenoviral
 CC vectors or clones which can be tested in vitro in an animal model.
 CC Furthermore, the methods produce RCA-free adenoviral libraries. RCA
 CC (replication competent adenovirus) contamination throughout the libraries
 CC could become a major obstacle, especially if libraries are continuously
 CC amplified for use in multiple screening programs. Additionally, a further
 CC advantage is minimisation of the number of steps involved in the process.
 CC There is no requirement for cloning each individual adenovirus before use
 CC in a high throughput screening program. Other systems include one or more
 CC steps in E. coli to achieve homologous recombination for the various
 CC adenoviral plasmids necessary for vector generation. The present
 CC sequence is a linker sequence used in the construction of the adenoviral
 CC vector library of the invention.
 XX
 XX Sequence 55 BP; 9 A; 17 C; 17 G; 12 T; 0 other;
 XX
 Query Match 92.7%; Score 51; DB 24; length 55;
 Best Local Similarity 100.0%; Pred. No. 3.e-09;
 Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 QY 5 ATTGACCTAGTGGCGCGGCAAGCCGGCGGCGGCACTAGTCAATGCAAT 55
 DB 55 ATTGACCTAGTGGCGCGGCGGCAAGCCGGCGGCGGCACTAGTCAATGCAAT 55
 RESULT 9
 AAT48644
 ID AAT48644 standard; DNA; 50 BP.

XX
 XX AAT48644;
 AC
 XX
 XX 21-MAY-1997 (first entry)
 DT
 XX
 XX Synthetic hairpin oligonucleotide HP/asp1.
 DE
 XX
 XX Gene therapy; vaccine; vector; adenovirus; packaging system;
 KM hairpin; PICL; ss.
 KW
 XX
 XX Synthetic.
 OS
 PN WO9700326-A1.
 XX
 PD 03-JAN-1997.
 XX
 PF 14-JUN-1996; 96WO-NL00244.
 XX
 PR 26-JUN-1995; 95EP-0201728.
 XX
 PR 15-JUN-1995; 95EP-0201611.
 XX
 PA (INTR-) INTRIGENE BV.
 PA (UYE-) RIKSUNNY LTDEN.
 XX
 PI Bout A, Pallaux FJ, Hoebein RC, Valerio D, Van Der EbbaJ;
 XX
 DR WPI; 1997-077531/07.
 XX
 PT New packaging cells and nucleic acids for recombinant adenovirus -
 PT have no overlapping sequences, prevents homologous recombination;
 PT for use in gene therapy and vaccination
 XX
 PS Disclosure; Page 55; 88pp; English.
 XX
 CC Synthetic oligonucleotides HP/asp1 (AAT48644) and HP/asp2 (AAT48645)
 CC were used to generate a synthetic hairpin, recreating an Asp718
 CC site at one of the termini if inserted in the Asp718 site of
 CC adenovirus minimal vector PICL (see also AAT48630). Insertion of
 CC the oligonucleotides into PICL generated clone PICLhac (correct
 CC orientation) and PICLhac (reverse, non-functional orientation).
 CC The constructs were used to demonstrate the competence of a
 CC synthetic DNA sequence, that is capable of forming a hairpin
 CC structure, to serve as a primer for reverse strand synthesis in
 CC the generation of double-stranded DNA molecules in cells that
 CC contain and express adenovirus genes.
 XX
 XX Sequence 50 BP; 10 A; 17 C; 17 G; 6 T; 0 other;
 XX
 Query Match 86.2%; Score 47.4; DB 18; length 50;
 Best Local Similarity 98.0%; Pred. No. 7.1e-08;
 Matches 48; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 XX
 QY 1 GTACATTGACCTAGTGGCGCGGCAAGCCGGCGGCGGCACTAGTCA 49
 DB 1 GTACATTGACCTAGTGGCGCGGCGGCAAGCCGGCGGCGGCACTAGTCA 49
 RESULT 10
 AAF30231
 ID AAF30231 standard; DNA; 50 BP.
 XX
 XX AAF30231;
 AC
 XX
 XX 30-APR-2001 (first entry)
 DT
 XX
 XX Oligonucleotide forming hairpin structure.
 DE
 XX
 XX Adenovirus; vector; gene therapy; packaging cell; hairpin; ds.
 KM
 XX
 XX Synthetic.
 OS
 PN
 XX
 XX Key Location/Qualifiers
 FH misc_feature 1..4
 FT

```

PT      /tag- a
PT      /note- "single-stranded 5' overhang"
PT      50
PT      misc-feature      /tag- b
PT      /note- "single-stranded overhang on complementary
PT      strand of sequence 5'-GTAC-3'"
XX      WO200105945-A2.
XX      25-JAN-2001.
XX      19-JUL-2000; 2000WO-BP07074.
XX      19-JUL-1999; 99US-0356575.
XX      (INTR-) INTROGENE BV.
XX      Hoeben RC, Bout A, Valerio D, Van Der Eb AJ, Schouten G;
XX      Fallaux FJ;
XX      WPI; 2001-14734/15.
XX      Producing recombinant adenovirus for use in gene therapy, comprises
XX      culturing cells containing adenoviral nucleic acid having an
XX      encapsidating signal and inverted terminal repeat, and lacking
XX      overlapping sequences -
XX      Example; Page 39; 97pp; English.
XX      The present sequence is that of an oligonucleotide formed from 2
XX      partially complementary oligonucleotides creating a hairpin
XX      structure. The oligonucleotide forms an Asp718 recognition site
XX      when inserted into the Asp718 site of plasmid pICL (see A930333).
XX      This was performed as part of an experiment to determine whether
XX      the hairpin could be used as a primer for reverse strand synthesis
XX      on the displaced strand after replication had started from the
XX      inverted terminal repeat (ITR) of the vector. In adenovirus
XX      infected cells, linear DNA fragments have on one terminus an
XX      adenovirus-derived ITR and at the other terminus a sequence that
XX      can anneal to the same strand, when present in single-stranded
XX      form, thereby generating a hairpin structure, and will be
XX      converted to structures with ITRs at both ends. The resulting DNA
XX      molecules will replicate by the same mechanism as the wild-type
XX      adenovirus genomes. The invention provides adenovirus vectors and
XX      packaging cell lines useful in the safe generation of EI-deleted
XX      recombinant adenovirus vectors for gene therapy applications.
XX      Packaging cells contain adenovirus nucleic acids having an
XX      encapsidating signal and ITR, but lack sequences that overlap with
XX      the vector, thereby preventing homologous recombination.
XX      Sequence 50 BP; 10 A; 17 C; 17 G; 6 T; 0 other;
XX      Query Match      86.2%; Score 47.4; DB 22; Length 50;
XX      Best Local Similarity 98.0%; Pred. No. 7,1e-08;
XX      Matches 48; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX      1 GTACACTGACCTAGTGGCCGCCGCGCAAGCCCGGCGGCACTAGTCA 49
XX      1 GTACACTGACCTAGTGGCCGCCGCGCAAGCCCGGCGGCACTAGTCA 49
XX      RESULT 11
XX      A9347030
XX      ID A9347030 standard; DNA: 50 BP.
XX      A9347030;
XX      05-JUN-2002 (first entry)
XX      Adenovirus vectro pICLnac/haw linker sequence hp/asp1#1.
XX      DE Adenovirus vector library; ss; linker; high throughput screening;
XX      KA RCA; replication competent adenovirus.

```

```

XX      Synthetic.
XX      US6340595-B1.
XX      22-JAN-2002.
XX      21-JUL-1999; 99US-0358036.
XX      12-JUN-1998; 98US-0097239.
XX      (GALA-) GALAPAGOS GENOMICS NV.
XX      Vogels R, Bout A, Van Es H, Schouten G;
XX      WPI; 2002-224926/28.
XX      Library of expressible nucleic acids, useful for determining nucleic
XX      acid function, comprises one or more adenoviral vectors capable of
XX      transfecting a host cell with the nucleic acid -
XX      Example 3; Column 84; 111pp; English.
XX      The invention relates to a library (I) of a multitude of unique
XX      expressible nucleic acids (NA), comprises a number of compartments
XX      (II), each consisting of one or more adenoviral vectors (III)
XX      comprising at least one unique NA of (I) in an aqueous medium, where
XX      (III) is capable of introducing the NA into a host cell (IV), is
XX      capable of expressing the product of the NA in (IV), and is deleted in
XX      a portion of the adenoviral genome necessary for replication. Also
XX      included is a method for producing the library. The library is useful for
XX      determining the function of at least one nucleic acid that is present.
XX      The library uses high throughput generation of recombinant adenoviral
XX      vector libraries containing one or more sample nucleic acids, followed by
XX      high throughput screening of the adenoviral vector libraries in a host to
XX      alter the phenotype of the host as a means of assigning a function to
XX      expression product(s) of the sample nucleic acids. The entire process
XX      lends itself to automation especially when implemented in a 96-well or
XX      other multi-well format. The high throughput screening, using a number of
XX      different in vitro assays, provides a means of efficiently obtaining
XX      functional information in a relatively short period of time. The
XX      member(s) of the recombinant adenoviral libraries that exhibit or induce
XX      a desired phenotype in a host in vitro or in situ are identified to
XX      reduce the libraries to a manageable number of recombinant adenoviral
XX      vectors or clones which can be tested in vitro in an animal model.
XX      Furthermore, the methods produce RCA-free adenoviral libraries. RCA
XX      (replication competent adenovirus) contamination throughout the libraries
XX      could become a major obstacle, especially if libraries are continuously
XX      amplified for use in multiple screening programs. Additionally, a further
XX      advantage is minimisation of the number of steps involved in the process.
XX      There is no requirement for cloning each individual adenovirus before use
XX      in a high throughput screening program. Other systems include one or more
XX      steps in E. coli to achieve homologous recombination for the various
XX      adenoviral plasmids necessary for vector generation. The present
XX      sequence is a linker sequence used in the construction of the adenoviral
XX      vector library of the invention.
XX      Sequence 50 BP; 10 A; 17 C; 17 G; 6 T; 0 other;
XX      Query Match      86.2%; Score 47.4; DB 24; Length 50;
XX      Best Local Similarity 98.0%; Pred. No. 7,1e-08;
XX      Matches 48; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX      1 GTACATGACCTAGTGGCCGCCGCGCAAGCCCGGCGGCACTAGTCA 49
XX      1 GTACATGACCTAGTGGCCGCCGCGCAAGCCCGGCGGCACTAGTCA 49
XX      Db      1 GTACATGACCTAGTGGCCGCCGCGCAAGCCCGGCGGCACTAGTCA 49
XX      RESULT 12
XX      AA259131
XX      ID AA259131 standard; DNA: 45 BP.
XX      AA259131;

```

XX	11-APR-2000	(first entry)
DF	Hairpin structure-forming oligonucleotide.	
XX	Expressible nucleic acid library; gene expression; gene function;	
DE	capillary formation; cell proliferation; hairpin structure; ss.	
KW	Synthetic.	
OS	WO9964587-A2.	
PN	16-DEC-1999.	
XX	11-JUN-1999.	99MO-NL00367.
XX	12-JUN-1998.	98US-0097239.
PR	(INTR-) INTROGENE BV.	
PA	Schouten G, Vogels R, Bout A, Van Es H;	
P1	WPI: 2000-097536/08.	
DR	New library of expressible nucleic acids, useful for high-throughput screening of gene function, especially for identifying therapeutic molecules -	
PT	Disclosure; Fig 15; 223pp; English.	
PS	The invention relates to a library of expressible nucleic acids (NA) which contains many compartments, each comprising at least one vehicle comprising at least one NA, the vehicle being capable of efficiently introducing a NA into a cell for expression. The library is useful for determining the function of one or more nucleic acids within the library, or to screen for an expressible nucleic acid with a particular desired function. It is especially useful for high throughput screening of gene function for functional genomics applications and for screening for nucleic acids with potential therapeutic value. Cell types appropriate for selection of a particular phenotype may be useful for capillary formation and cell proliferation. This oligonucleotides is used to generate a hairpin structure which was used to determine if it could be used to prime reverse strand synthesis on the displaced strand after replication initiation in the adenoviral inverted terminal repeat (ITR).	
CC	Sequence 45 BP; 9 A; 16 C; 15 G; 5 T; 0 other:	
CC	Query Match	78.9%; Score 43.4; DB 21; Length 45;
CC	Best Local Similarity	97.8%; Pred. No. 1.9e-06;
CC	Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	
OY	1 GTACATGACCTAGTGGCGGCCGGGAAMGCCGGGGGCGCACTAG 45	
DB		
	1 GTACACTGACCTTAGTGCCGCCCGGCACAAKCGCGCGGCGCACTAG 45	
RESULT 13		
AAZ37961		
ID	AAZ37961 standard; DNA; 45 BP.	
XX	AAZ37961;	
XX	07-FEB-2000 (first entry)	
XX	DNA molecule containing Rp/asp sequences.	
DE	Adenoviral vector; replication-defective; adenovirus; ITR; hepatitis;	
KW	inverted terminal repeat; encapsulation signal; gene therapy; tumor;	
KW	hypercholesterolemia; cystic fibrosis; Duchenne muscular dystrophy;	
KW	autoimmune disease; rheumatoid arthritis; ss.	
XX	Synthetic.	

```

XX      MO9955132-A2.
PX      04-NOV-1999.
XX      23-APR-1999;      99MO-NL00235.
XX      24-APR-1998;      98US-0065752.
XX      (INTR-) INTRIGENE BV.
PA      (INTR-) INTRIGENE BV.
XX      Vogels R, Bout A;
PI      MPI; 2000-023229/02.
XX      New recombinant adenovirus vectors, used particularly for gene therapy
PT      for treating inherited or acquired diseases
XX      Disclosure; Fig 15; 16pp; English.
PS      The invention provides methods of producing recombinant adenoviral
CC      vectors (Adv's) for generating replication-defective adenoviruses.
CC      Generating an Adv comprises fusing 2 partially overlapping sequences
CC      nucleic acid molecules that are capable of combining with each other to
CC      allow the generation of a physically linked nucleic acid comprising at
CC      least 2 functional adenoviral inverted terminal repeats (ITRs), a
CC      functional encapsulation signal and a nucleic acid of interest. The
CC      products can be used for gene therapy for treating inherited diseases
CC      e.g. cystic fibrosis, Duchenne molecular dystrophy, hypercholesterolemia,
CC      blood clotting disorders (hemophilia) or acquired diseases such as
CC      tumors, hepatitis, (auto)immune diseases, restenosis, or Rheumatoid
CC      arthritis.
XX      Sequence 45 BP; 9 A; 16 C; 15 G; 5 T; 0 other:
SQ
XX      Query Match      78.9%; Score 43.4; DB 21; Length 45;
XX      Best Local Similarity 97.8%; Pred. No. 1.9e-06;
XX      Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY      1 GTACATTGACCTAGTGCCTCCGCGGCAAGCCCGGCGGACTAG 45
      1 ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB      1 GTACACTGACCTAGTGCCTCCGCGGCAAGCCCGGCGGACTAG 45
XX
XX      RESULT 14
XX      AAF30234
XX      ID AAF30234 standard; DNA; 45 BP.
XX      AAF30234;
XX      AC AAF30234;
XX      DT 30-APR-2001 (first entry)
XX      DE Oligonucleotide forming hairpin structure.
XX      YY
XX      KW Adenovirus; vector; gene therapy; packaging cell; hairpin; ss.
XX      OS Synthetic.
XX      PN WO200105945-A2.
XX      PP 25-JAN-2001.
XX      PF 19-JUL-2000; 2000MO-EP07074.
XX      PR 19-JUL-1999; 99US-0356575.
XX      PA (INTR-) INTRIGENE BV.
XX      PI Hoeben RG, Bout A, Valerio D, Van Der Eb AJ, Schouten G;
XX      Fallaux FG;
XX      MPI; 2001-147334/15.
XX

```

PT Producing recombinant adenovirus for use in gene therapy, comprises
 PT culturing cells containing adenoviral nucleic acid having an
 PT encapsulating signal and inverted terminal repeat, and lacking
 PT overlapping sequences -
 XX
 XX
 XX Example; Page 53; 97pp; English.

XX The present sequence is that of an oligonucleotide that forms a
 CC hairpin structure. The oligonucleotide creates an Asp718 recognition
 CC site when inserted into the Asp718 site of plasmid pICL (see
 CC AAF30233). This was performed as part of an experiment to determine
 CC whether the hairpin could be used as a primer for reverse strand
 CC synthesis on the displaced strand after replication had started from
 CC the inverted terminal repeat (ITR) of the vector. In adenovirus
 CC infected cells, linear DNA fragments have on one terminus an
 CC adenovirus-derived ITR and at the other terminus a sequence that
 CC can anneal to the same strand, when present in single-stranded
 CC form, thereby generating a hairpin structure, and will be
 CC converted to structures with ITRs at both ends. The resulting DNA
 CC molecules will replicate by the same mechanism as the wild-type
 CC adenovirus genomes. The invention provides adenovirus vectors and
 CC packaging cell lines useful in the safe generation of EI-deleted
 CC recombinant adenovirus vectors for gene therapy applications.
 CC Packaging cells contain adenovirus nucleic acids having an
 CC encapsulating signal and ITR, but lack sequences that overlap with
 CC the vector, thereby preventing homologous recombination.
 CC
 XX

SO Sequence 45 BP; 9 A; 16 C; 15 G; 5 T; 0 other;

Query Match 78.9%; Score 43.4; DB 22; Length 45;
 Best Local Similarity 97.8%; Pred. No. 1.9e-06;
 Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GTACATTGACCTAGTCCGCCCGGCAAGCCCGGCGGCACTAG 45
 DB 1 GTACACTGACTAGTCCGCCCGGCAAGCCCGGCGGCACTAG 45

RESULT 15
 ABK47038 ID ABK47038 standard; DNA: 45 BP.
 XX
 XX AC ABK47038;
 XX
 XX DT 05-JUN-2002 (first entry)
 XX
 XX DE Adenovirus vector potential hairpin forming sequence.
 XX
 XX KM Adenovirus vector library; ss: high throughput screening; Asp7181;
 XX RCA: replication competent adenovirus; hairpin; pICLHa.
 OS Mastadenovirus Ad5.
 XX
 XX US6340595-B1.
 XX
 XX PN 22-JAN-2002.
 XX
 XX PD 21-JUN-1999; 99US-0358036.
 XX
 XX PF 12-JUN-1998; 98US-0097239.
 XX
 XX PR (GALA-) GALAPAGOS GENOMICS NV.
 XX
 XX PA Vogels R, Rout A, Van Es H, Schouten G;
 XX
 XX PI WPI: 2002-224926/28.
 XX
 XX DR

PT Library of expressible nucleic acids, useful for determining nucleic
 PT acid function, comprises one or more adenoviral vectors capable of
 PT transfecting a host cell with the nucleic acid -
 XX
 XX Example 3; Fig 15; 111pp; English.

CC The invention relates to a library (I) of a multitude of unique
 CC expressible nucleic acids (NA), comprises a number of compartments
 CC (II), each consisting of one or more adenoviral vectors (III)
 CC comprising at least one unique NA of (I) in an aqueous medium, where
 CC (III) is capable of introducing the NA into a host cell (IV), is
 CC capable of expressing the product of the NA in (IV), and is deleted in
 CC a portion of the adenoviral genome necessary for replication. Also
 CC included is a method for producing the library. The library is useful for
 CC determining the function of at least one nucleic acid that is present.
 CC The library uses high throughput generation of recombinant adenoviral
 CC vector libraries containing one or more sample nucleic acids, followed by
 CC high throughput screening of the adenoviral vector libraries in a host to
 CC alter the phenotype of the host as a means of assigning a function to
 CC expression product(s) of the sample nucleic acids. The entire process
 CC lends itself to automation especially when implemented in a 96-well or
 CC other multi-well format. The high throughput screening, using a number of
 CC different *in vitro* assays, provides a means of efficiently obtaining
 CC functional information in a relatively short period of time. The
 CC member(s) of the recombinant adenoviral libraries that exhibit or induce
 CC a desired phenotype in a host *in vitro* or *in situ* are identified to
 CC reduce the libraries to a manageable number of recombinant adenoviral
 CC vectors or clones which can be tested *in vitro* in an animal model.
 CC Furthermore, the methods produce RCA-free adenoviral libraries. RCA
 CC (replication competent adenovirus) contamination throughout the libraries
 CC could become a major obstacle, especially if libraries are continuously
 CC amplified for use in multiple screening programs. Additionally, a further
 CC advantage is minimization of the number of steps involved in the process.
 CC There is no requirement for cloning each individual adenovirus before use
 CC in a high throughput screening program. Other systems include one or more
 CC steps in *E. coli* to achieve homologous recombination for the various
 CC adenoviral plasmids necessary for vector generation. The present
 CC sequence is part of the adenovirus vector pICLHa which can form a
 CC hairpin sequence on digestion with restriction endonuclease Asp7181.
 CC During chain elongation, the free 3' terminus becomes displaced and can
 CC act as a double-stranded template for cellular or viral DNA polymerase.
 CC
 XX

SO Sequence 45 BP; 9 A; 16 C; 15 G; 5 T; 0 other;

Query Match 78.9%; Score 43.4; DB 24; Length 45;
 Best Local Similarity 97.8%; Pred. No. 1.9e-06;
 Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GTACATTGACCTAGTCCGCCCGGCAAGCCCGGCGGCACTAG 45
 DB 1 GTACACTGACTAGTCCGCCCGGCAAGCCCGGCGGCACTAG 45

Search completed: December 27, 2002, 04:46:15
 Job time : 172.5 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 Comugen Ltd.

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

OM nucleic acid - nucleic acid search, using SW model

Run on: December 27, 2002, 03:13:39 ; Search time 1330 seconds

(without alignments)
1203.499 Million cell updates/sec

Title: US-09-918-029-19

Perfect score: 55
Sequence: 1 ggcacgtacactagtcgcgc.....ggcgacactagtcacatgat 55

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

Genbank: *
1: gb_da: *
2: gb_htg: *
3: gb_in: *
4: gb_om: *
5: gb_ov: *
6: gb_pat: *
7: gb_ph: *
8: gb_pl: *
9: gb_pt: *
10: gb_ro: *
11: gb_sts: *
12: gb_sy: *
13: gb_un: *
14: gb_vl: *
15: em_da: *
16: em_fun: *
17: em_hum: *
18: em_in: *
19: em_mu: *
20: em_om: *
21: em_ov: *
22: em_ph: *
23: em_pat: *
24: em_pl: *
25: em_ro: *
26: em_sts: *
27: em_un: *
28: em_vl: *
29: em_hum: *
30: em_hum: *
31: em_hum: *
32: em_hum: *
33: em_hum: *
34: em_hum: *
35: em_hum: *
36: em_hum: *
37: em_hum: *
38: em_hum: *
39: em_hum: *
40: em_hum: *
41: em_hum: *
42: em_hum: *
43: em_hum: *
44: em_hum: *
45: em_hum: *

Pred. No. is the number of results predicted by chance to have a

Result No.	Score	Query Match	Length	DB ID	Description
1	55	100.0	55	6	AR091516
2	55	100.0	55	6	AR154396
3	55	100.0	55	6	AR174324
4	55	100.0	55	6	AR183308
5	55	100.0	55	6	AX080360
6	55	92.7	55	6	AR091517
7	55	92.7	55	6	AR154397
8	55	92.7	55	6	AR174325
9	55	92.7	55	6	AR183309
10	55	92.7	55	6	AX080361
11	47.4	86.2	50	6	AR091514
12	47.4	86.2	50	6	AR154394
13	47.4	86.2	50	6	AR174322
14	47.4	86.2	50	6	AR183306
15	47.4	86.2	50	6	AX080358
16	43.4	78.9	45	6	AR091519
17	43.4	78.9	45	6	AR154399
18	43.4	78.9	45	6	AR174327
19	43.4	78.9	45	6	AR183314
20	43.4	78.9	50	6	AR091515
21	43.4	78.9	50	6	AR154395
22	43.4	78.9	50	6	AR174323
23	43.4	78.9	50	6	AR183307
24	43.4	78.9	50	6	AX080359
25	42.2	76.7	55	6	AR091516
26	42.2	76.7	55	6	AR091517
27	42.2	76.7	55	6	AR154396
28	42.2	76.7	55	6	AR174324
29	42.2	76.7	55	6	AR183315
30	42.2	76.7	55	6	AX080362
31	42.2	76.7	55	6	AR183308
32	42.2	76.7	55	6	AR183309
33	42.2	76.7	55	6	AX080360
34	42.2	76.7	55	6	AX080361
35	40.2	73.1	45	6	AX080363
36	38.6	70.2	50	6	AR091514
37	38.6	70.2	50	6	AR091515
38	38.6	70.2	50	6	AR154394
39	38.6	70.2	50	6	AR154395
40	38.6	70.2	50	6	AR174322
41	38.6	70.2	50	6	AR174323
42	38.6	70.2	50	6	AR183306
43	38.6	70.2	50	6	AR183307
44	38.6	70.2	50	6	AX080358
45	38.6	70.2	50	6	AX080359

ALIGNMENTS

RESULT 1
AR091516
LOCUS AR091516
DEFINITION Sequence 19 from patent US 5994128.
ACCESSION AR091516
VERSION AR091516.1 GI:10018271
KEYWORDS
SOURCE
ORGANISM
Unknown.
REFERENCE
1 (bases 1 to 55)
AUTHORS Fallaux F, Jacobus, Hoebe, R, Cornelis, Van der Ed, A, Jan., Bout, A.
and Valerio, D.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy

Pred. No. is the number of results predicted by chance to have a

KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 55)
AUTHORS Fallaux,F.Jacobus., Hoeben,R.Cornelis., Van der Eb,A.Jan., Bout,A. and Valerio,D.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL Patent: US 5994128-A 20 30-NOV-1999;
FEATURES Location/Qualifiers
source 1..55
/organism="unknown"
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN
Query Match 92.7%; Score 51; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 6.2e-07;
Matches 51: Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 5 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 55
Db 55 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 5
RESULT 7
AR154397/c AR154397 55 bp DNA linear PAT 08-AUG-2001
LOCUS Sequence 20 from patent US 6238893.
DEFINITION AR154397
ACCESSION AR154397
VERSION AR154397.1 GI:15122450
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 55)
AUTHORS Hoeben,R.Cornelis. and Bout,A.
TITLE Method for intracellular DNA amplification
JOURNAL Patent: US 6238893-A 20 29-MAY-2001;
FEATURES Location/Qualifiers
source 1..55
/organism="unknown"
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN
Query Match 92.7%; Score 51; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 6.2e-07;
Matches 51: Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 5 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 55
Db 55 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 5
RESULT 8
AR174325/c AR174325 55 bp DNA linear PAT 17-DEC-2001
LOCUS Sequence 20 from patent US 6306552.
DEFINITION AR174325
ACCESSION AR174325
VERSION AR174325.1 GI:17914645
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 55)
AUTHORS Fallaux,F.Jacobus., Hoeben,R.Cornelis., Van der Eb,A.Jan., Bout,A. and Valerio,D.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL Patent: US 6306552-A 20 23-OCT-2001;
FEATURES Location/Qualifiers
source 1..55
/organism="unknown"

BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN
Query Match 92.7%; Score 51; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 6.2e-07;
Matches 51: Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 5 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 55
Db 55 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 5
RESULT 9
AR183309/c AR183309 55 bp DNA linear PAT 20-APR-2002
LOCUS Sequence 42 from patent US 6340595.
DEFINITION AR183309
ACCESSION AR183309
VERSION AR183309.1 GI:20226902
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 55)
AUTHORS Vogels,R., Bout,A., van Es,H. and Schouten,G.
TITLE High throughput screening of gene function using adenoviral libraries for functional genomics applications
JOURNAL Patent: US 6340595-A 42 22-JAN-2002;
FEATURES Location/Qualifiers
source 1..55
/organism="unknown"
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN
Query Match 92.7%; Score 51; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 6.2e-07;
Matches 51: Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 5 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 55
Db 55 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 5
RESULT 10
AX080361/c AX080361 55 bp DNA linear PAT 22-FEB-2001
LOCUS Sequence 20 from Patent WO0105945.
DEFINITION AX080361
ACCESSION AX080361
VERSION AX080361.1 GI:13159819
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 55)
AUTHORS Hoeben,R.C., Bout,A., Valerio,D., van der Eb,A.J., Schouten,G. and Fallaux,F.J.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL Patent: WO 0105945-A 20 25-JAN-2001;
FEATURES Introgene B.V. (NL)
source 1..55
Location/Qualifiers
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="oligonucleotide HP/c1a2"
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN
Query Match 92.7%; Score 51; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 6.2e-07;
Matches 51: Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 5 ATTGACCTAGTGGCCCGGCAAAAGCCGGCGGCACTAGTCAATCGAT 55
|||||

Db	55	ATTCACCTAGTCCGCCCGGGCAAAAGCCCGGGCGGCACTAGTCAATGCAT	5
RESULT	11		
LOCUS	AR091514	50 bp	DNA
DEFINITION	Sequence 17 from patent US 5994128.		Linear
ACCESSION	AR091514		
VERSION	AR091514.1	GI:10018269	
KEYWORDS			
SOURCE	Unknown.		
ORGANISM	Unknown.		
REFERENCE	1 (bases 1 to 50)		
AUTHORS	Fallaix,F.Jacobus., Hoeben,R.Cornelis., Van der Ed,A.Jan., Bout,A. and Valerio,D.		
TITLE	Packaging Systems for human recombinant adenovirus to be used in gene therapy		
JOURNAL	Patent: US 5994128-A 17 30-NOV-1999;		
FEATURES	Location/Qualifiers		
source	1..50		
BASE COUNT	10 a 17 c 17 g 6 t		
ORIGIN	/organism="unknown"		
Query Match	86.2%; Score 47.4; DB 6; Length 50;		
Best Local Similarity	98.0%; Pred. No. 1e-05;		
Matches	48; Conservative 0; Mismatches 1; Indels 0; Gaps 0;		
Oy	1 GTACATTGACCTAGTCCGCCCGGGCAAAAGCCCGGGCGGCACTAGTCA 49		
Db	1 GTACACTGACCTAGTCCGCCCGGGCAAAAGCCCGGGCGGCACTAGTCA 49		
RESULT	12		
LOCUS	ARI54394	50 bp	DNA
DEFINITION	Sequence 17 from patent US 6238893.		Linear
ACCESSION	ARI54394		
VERSION	ARI54394.1	GI:15122447	
KEYWORDS			
SOURCE	Unknown.		
ORGANISM	Unknown.		
REFERENCE	1 (bases 1 to 50)		
AUTHORS	Hoeben,R.Cornelis. and Bout,A.		
TITLE	Method for intracellular DNA amplification		
JOURNAL	Patent: US 6238893-A 17 29-MAY-2001;		
FEATURES	Location/Qualifiers		
source	1..50		
BASE COUNT	10 a 17 c 17 g 6 t		
ORIGIN	/organism="unknown"		
Query Match	86.2%; Score 47.4; DB 6; Length 50;		
Best Local Similarity	98.0%; Pred. No. 1e-05;		
Matches	48; Conservative 0; Mismatches 1; Indels 0; Gaps 0;		
Oy	1 GTACATTGACCTAGTCCGCCCGGGCAAAAGCCCGGGCGGCACTAGTCA 49		
Db	1 GTACACTGACCTAGTCCGCCCGGGCAAAAGCCCGGGCGGCACTAGTCA 49		
RESULT	13		
LOCUS	ARI174322	50 bp	DNA
DEFINITION	Sequence 17 from patent US 630652.		Linear
ACCESSION	ARI174322		
VERSION	ARI174322.1	GI:17914642	
KEYWORDS			
SOURCE	Unknown.		
ORGANISM	Unknown.		
	Unclassified.		

REFERENCE	1 (bases 1 to 50)			
AUTHORS	Fallaux,F.Jacobus., Hoeben,R.Cornelius., Van Der Eb,A.Jan., Bout,A. and Valerio,D.			
TITLE	Packaging systems for human recombinant adenovirus to be used in gene therapy			
JOURNAL	Patent: US 630652-A 17 23-OCT-2001;			
FEATURES	Location/Qualifiers			
source	1..50			
BASE COUNT	10 a 17 c 17 g 6 t			
ORIGIN				
Query Match	86.2%; Score 47.4; DB 6; Length 50;			
Best Local Similarity	98.0%; Pred.No. 1e-05;			
Matches	48; Conservative 0; Mismatches 1; Indels 0; Gaps 0;			
QY	1 GTACCTGACCTAGTGGCGCGGCAAAAGCCGGGCGGCACTAGGTCA 49			
Db	1 GTACCTGACCTAGTGGCGCGGCAAAAGCCGGGCGGCACTAGGTCA 49			
RESULT 14				
LOCUS	ARI83306 50 bp DNA linear PAT 20-APR-2002			
DEFINITION	Sequence 39 from patent US 6340595.			
ACCESSION	ARI83306			
VERSION	ARI83306.1 GI:20226899			
KEYWORDS	.			
SOURCE	Unknown.			
ORGANISM	Unknown.			
REFERENCE	Unclassified.			
AUTHORS	1 (bases 1 to 50)			
TITLE	Vogels,R., Bout,A., van Es,H. and Schouten,G.			
JOURNAL	High throughput screening of gene function using adenoviral libraries for functional genomics applications			
FEATURES	Patent: US 6340595-A 39 22-JAN-2002;			
source	Location/Qualifiers			
	1..50			
BASE COUNT	10 a 17 c 17 g 6 t			
ORIGIN	/organism="unknown"			
Query Match	86.2%; Score 47.4; DB 6; Length 50;			
Best Local Similarity	98.0%; Pred.No. 1e-05;			
Matches	48; Conservative 0; Mismatches 1; Indels 0; Gaps 0;			
QY	1 GTACCTGACCTAGTGGCGCGGCAAAAGCCGGGCGGCACTAGGTCA 49			
Db	1 GTACCTGACCTAGTGGCGCGGCAAAAGCCGGGCGGCACTAGGTCA 49			
RESULT 15				
LOCUS	AX080358 50 bp DNA linear PAT 22-FEB-2001			
DEFINITION	Sequence 17 from Patent WO0105945.			
ACCESSION	AX080358			
VERSION	AX080358.1 GI:13159816			
KEYWORDS	.			
SOURCE	synthetic construct.			
ORGANISM	synthetic construct			
REFERENCE	artificial sequences.			
AUTHORS	1 (bases 1 to 50)			
TITLE	Hoeben,R.C., Bout,A., Valerio,D., van der Eb,A.J., Schouten,G. and Fallaux,F.J.			
JOURNAL	Packaging systems for human recombinant adenovirus to be used in gene therapy			
FEATURES	Patent: WO 0105945-A 17 25-JAN-2001;			
source	Introgene B.V. (NL)			
	Location/Qualifiers			
	1..50			
	/organism="synthetic construct"			
	/db_xref="taxon:32630"			
	/note="Oligonucleotide HP/aspl"			

BASE COUNT 10 a 17 c 17 g 6 t
ORIGIN

Query Match 86.2%; Score 47.4; DB 6; Length 50;
Best Local Similarity 98.0%; Pred. No. 1e-05;
Matches 48; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GTACATTGACCTAGTCCCGGCGCAAGCCGGGCGGCACTAGGTCA 49
|||||
Db 1 GTACACTGACCTAGTCCCGGCGCAAGCCGGGCGGCACTAGGTCA 49

Search completed: December 27, 2002, 05:31:02
Job time : 1333.5 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 27, 2002, 04:38:01 ; Search time 1321.5 Seconds
(without alignments)
674.046 Million cell updates/sec

Title: US-09-918-029-19

Perfect score: 55
1 gtcactgactagtcgcgc.....gcgcactagtcacatcat 55

Sequence: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Scoring table: Gapop 10.0, Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues 32308132

Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database :

EST:*

1: em_estba:*
2: em_esthum:*
3: em_estlin:*
4: em_estlmu:*
5: em_estlow:*
6: em_estlpl:*
7: em_estlro:*
8: em_hlc:*
9: gb_estl1:*
10: gb_estl2:*
11: gb_hlc:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estlin:*
16: em_estlmu:*
17: gb_gss:*
18: em_gss_hum:*
19: em_gss_lnv:*
20: em_gss_pln:*
21: em_gss_vrl:*
22: em_gss_fun:*
23: em_gss_mam:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rtd:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	25.2	45.8	794	10	BE294341 601172854
2	24.8	45.1	756	10	BE254363 601109113
3	24.6	44.7	749	12	BE254363 601109113
4	24.4	43.6	885	9	AL564683 102400761
5	23.8	43.3	174	17	AL564683 102400761
6	23.8	43.3	885	9	AL564683 102400761

C	7	23.6	42.9	1115	13	BM546853	BM546853	AGENCOURT
C	8	23.6	42.9	1208	12	BG474535	BG474535	602577304
C	9	23.4	42.5	596	12	BG194272	BG194272	RST13417
C	10	23.4	42.5	884	14	BQ892955	BQ892955	AGENCOURT
C	11	23.4	42.5	1149	14	BQ642928	BQ642928	AGENCOURT
C	12	23.2	42.2	344	9	AA081549	AA081549	zr21907.r
C	13	23.2	42.2	590	17	A2205863	A2205863	SP_0105.A
C	14	23.2	42.2	633	13	B1548352	B1548352	603189633
C	15	23.2	42.2	682	13	BE69165	BE69165	603295596
C	16	23.2	42.2	703	10	BE381912	BE381912	601272452
C	17	23.2	42.2	749	12	BG722376	BG722376	602693574
C	18	23.2	42.2	755	12	BG719715	BG719715	602690436
C	19	23.2	42.2	814	12	BG717933	BG717933	602633988
C	20	23.2	42.2	816	13	B1464359	B1464359	603204058
C	21	23.2	42.2	901	12	BG470218	BG470218	602533732
C	22	23.2	42.2	1049	14	BQ068336	BQ068336	AGENCOURT
C	23	23.2	42.2	1135	12	BG536831	BG536831	602566369
C	24	23.2	42.2	1235	14	BQ225342	BQ225342	AGENCOURT
C	25	23.2	41.8	530	13	B1727969	B1727969	103109550
C	26	23.2	41.8	597	14	BQ818268	BQ818268	103006961
C	27	23.2	41.8	607	13	B1726556	B1726556	103108640
C	28	23.2	41.8	670	14	BQ819135	BQ819135	103007560
C	29	23.2	41.8	708	14	BQ813738	BQ813738	1030038F0
C	30	23.2	41.8	714	12	BG848999	BG848999	1024023E1
C	31	23.2	41.8	720	12	BG849000	BG849000	1024023E1
C	32	23.2	41.8	820	13	BG923596	BG923596	602833455
C	33	23.2	41.8	822	17	CNS0211X	AL117646	Telradon
C	34	23.2	41.8	906	17	CNS03A5J	AL234784	Telradon
C	35	23.2	41.8	977	14	BQ672537	BQ672537	AGENCOURT
C	36	23.2	41.8	1030	14	BQ051246	BQ051246	AGENCOURT
C	37	22.8	41.5	319	17	AQ357767	AQ357767	CITR1-E1-
C	38	22.8	41.5	344	9	AA081549	AA081549	zr21907.r
C	39	22.8	41.5	526	12	BQ812739	BQ812739	1030032A0
C	40	22.8	41.5	565	12	BE725257	BE725257	894082A06
C	41	22.8	41.5	592	9	AL703582	DKF2P686C	AL703582
C	42	22.8	41.5	599	17	AQ095699	RPCI-23-3	AQ095699
C	43	22.8	41.5	658	12	BG724427	BG724427	602693726
C	44	22.8	41.5	749	12	BG722376	BG722376	602693574
C	45	22.8	41.5	755	12	BG719715	BG719715	602690436

ALIGNMENTS

RESULT 1
LOCUS BE294341 794 bp mRNA linear EST 20-JUL-2000
DEFINITION 601172854F1 NIH_MGC_17 Homo sapiens CDNA clone IMAGE:3528349 5',
mRNA sequence.
ACCESSION BE294341
VERSION BE294341.1 GI:9177788
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS NIH-MGC http://mgc.nci.nih.gov/
TITLE 1 (bases 1 to 794)
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cga@bbs.fmail.nih.gov

Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)
DNA Sequencing by: Incyte Genomics, Inc.
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNLN at: image.lnl.gov
Plate: L10K197 row: 1 column: 14
High quality sequence stop: 605.
Location/Qualifiers
1..794
/organism="Homo sapiens"

FEATURES

source

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE I (bases 1 to 885)
Li,W.B., Gruber,C., Jesse,J. and Polayes,D.
Full-length cDNA libraries and normalization
Unpublished (2001)

JOURNAL COMMENT Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr,
Location/Qualifiers

FEATURES
source
1..885
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="CSDDM007KK12"
/clone_1lb="LTI_NFLD01_NBRC4"
/sex="male"
/tissue_type="neuroblastoma cells"
/lab_host="DH10B"
/note="Organ: brain; Vector: pCMVSPORT 6; 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-stranded cDNA was digested with Not I and cloned into the Not I and Eco RV sites of the pCMVSPORT 6 vector. Library was normalized. Library was constructed by Life Technologies. Contact : Feng Liang Life Technologies, a division of Invitrogen 9800 Medical Center Drive Rockville, Maryland 20850, USA Fax : (1) 301 610 8371 Email : fliang@lifestech.com URL : http://fulllength.invitrogen.com"

BASE COUNT 166 a 253 c 263 g 159 t 44 others

ORIGIN

Query Match 43.6%; Score 24; DB 9; Length 885;
Best Local Similarity 63.5%; Pred. No. 4.7e+02;
Matches 33; Conservative 2; Mismatches 17; Indels 0; Gaps 0;

Oy 1 GTACATTGACTGATGCCTGGCCCGGCAAAAGCCCGGGCGGCACTAGCTCAATC 52
| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 612 GNAAGTGCGMGCSMTGTGTGCCGCGACTCAGCACCTCGCGCACTTTGTCAAGC 561

RESULT 5 AOS688337 174 bp DNA linear GSS 07-JUN-1999
AOS688337 LOCUS CITBI-EI-2644P17.TF CITBI-EI Homo sapiens genomic clone 2644P17,
DEFINITION DNA sequence.
ACCESSION AOS688337
VERSION AOS688337.1 GI:5015017
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 174)
Zhao,S., Adams,M.D., Niernan,W., Malek,J., Shizuya,H., Simon,M. and
Venier,J.C
Use of BAC End Sequences from Caltech Libraries for Sequence-Ready
Map Building
Unpublished (1997)
Other_GSSs: CITBI-EI-2644P17.TF
Contact: Shaying Zhao, William Niernan, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbeetjgr.org
Clones are available from Research Genetics (info@resgen.com). BAC
end search page:
http://www.tigr.org/tdb/hungen/Bac_end_search/bac_end_search.html.
Seq primer: M3 Reverse
Class: BAC ends.

FEATURES		Location/Qualifiers	
source		1..174	
		/organism="Homo sapiens"	
		/db_xref="taxon.9606"	
		/clone="2644p17"	
		/clone_1b="C17H1-E1"	
		/sex="male"	
		/cell_type="sperm"	
		/note="Vector: pHELOBAC11; Site_1: EcoRI; Site_2: EcoRI; CalTech Human BAC Library D"	
BASE COUNT		47 a	47 c 49 g 30 t 1 others
ORIGIN			
Query Match		43.3%	Score 23.8; DB 17; Length 174;
Best local Similarity		72.1%;	Pred. No. 3.6e+02;
Matches 31; Conservative		0;	Mismatches 12; Indels 0; Gaps 0;
OY	8 GACCTAGTCCGCCGCAAAAGCCCGGCGCACTAGTCATC 50		
Db	118 GACCTTCGTCTCCGGGAAAGCCAGTAGTACTAAGCA 76		
RESULT 6		885 bp	mRNA linear EST 16-FEB-2001
AL564683			
LOCUS			
DEFINITION		AL564683 LTI_NFL001_NBC4 Homo sapiens CDNA clone CS0DH007K12 3	
ACCESSION		prime, mRNA sequence.	
VERSION		AL564683	
KEYWORDS		AL564683.1 GI:12915335	
SOURCE		EST.	
ORGANISM		human.	
REFERENCE		Homo sapiens	
AUTHORS		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
TITLE		Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.	
JOURNAL		1 (bases 1 to 885)	
COMMENT		Li, W.B., Gruber, C., Jesse, J. and Polayes, D.	
		Full-length CDNA libraries and normalization	
		Unpublished (2001)	
		Contact: Genoscope	
		Genoscope - Centre National de Sequencage	
		BP 191 91006 EVRY cedex - France	
		Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.	
FEATURES		Location/Qualifiers	
source		1..885	
		/organism="Homo sapiens"	
		/db_xref="taxon.9606"	
		/clone="CS0DH007K12"	
		/clone_1b="LTI_NFL001_NBC4"	
		/sex="male"	
		/tissue_type="neuroblastoma cells"	
		/lab_host="DH10B"	
		/note="Organ: brain; Vector: PCMVSPORT 6; 1st strand CDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-stranded CDNA was digested with Not I and cloned into the Not I and Eco RV sites of the PCMVSPORT 6 vector. Library was normalized. Library was constructed by life technologies. Contact : Feng Liang Life Technologies, a division of invitrogen 9800 Medical Center Drive Rockville, Maryland 20850, USA Fax : (1) 301 610 8371 Email : fliang@lifetech.com URL : http://fulllength.invitrogen.com"	
BASE COUNT		166 a 253 c 263 g 159 t 44 others	
ORIGIN			
Query Match		43.3%	Score 23.8; DB 9; Length 885;
Best local Similarity		66.0%;	Pred. No. 5.5e+02;
Matches 31; Conservative		2;	Mismatches 16; Indels 0; Gaps 0;
OY	6 TTGACCTAGTCCGCCGCAAAAGCCCGGCGCACTAGTCATC 52		
Db	563 TTGACACAGTCCGCGAGTGTGTGAGCTCGCGGACACACGCKCACCC 609		

REFERENCE	LOCUS	DEFINITION	ACCESSION	VERSION	KEYWORDS	SOURCE	ORGANISM
BM546853/7	BM546853	1115 bp	MRNA	linear	EST 20-FEB-2002		
LOCUS	AGNCOURT	6491290	NIH_MGC_125	Homo sapiens	CDNA clone IMAGE:5723591		
DEFINITION	5', MRNA sequence.						
ACCESSION	BM546853						
VERSION	BM546853.1	GI:18780144					
KEYWORDS	EST.						
SOURCE	human.						
ORGANISM	Homo sapiens						
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;						
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.						
TITLE	NIH-MGC http://mhc.nci.nih.gov/.						
JOURNAL	National Institutes of Health, Mammalian Gene Collection (MGC)						
COMMENT	Unpublished (1999)						
	Contact: Robert Strausberg, Ph.D.						
	Email: cgraphs-remail.nih.gov						
	Tissue Procurement: Invitrogen						
	CDNA Library Preparation: Life Technologies, Inc.						
	CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)						
	DNA Sequencing by: Agencourt Bioscience Corporation						
	Clone distribution: MGC clone distribution information can be						
	found through the I.M.A.G.E. Consortium/LNL at:						
	http://image.llnl.gov						
	Plate: L14M12711 row: h column: 24						
	High quality sequence start: 102						
	High quality sequence stop: 420.						
FEATURES	Location/Qualifiers						
SOURCE	1..1115						
	/organism="Homo sapiens"						
	/db_xref="taxon:9606"						
	/clone="IMAGE:5723591"						
	/clone_1ib="NIH_MGC_125"						
	/lab_host="DH10B"						
	/note="organ: ovary (pool of 3); Vector: pCMV-SPORT6;						
	site.1: EcoRV (destroyed); site.2: NotI; RNA source pool						
	of three ovaries, from females ranging in age from 38 to						
	49 yo. Library is oligo-dT primed and directionally cloned						
	(EcoRV site is destroyed upon cloning). Average insert						
	size 2.1 kb, insert size range 1-3.5 kb. Library is						
	normalized and enriched for full-length clones and was						
	constructed by C. Gruber (Invitrogen). Research Genetics						
	tracking code 036."						
BASE COUNT	196 a 365 c 343 g 208 t 3 others						
ORIGIN							
	Query Match	42.9%; Score 23.6; DB 13; Length 1115;					
	Best local Similarity 69.6%; Pred. No. 6.8e+02;						
	Matches 32; Conservative 0; Mismatch 14; Indels 0; Gaps 0;						
Qy	4 CATTGACCTAGTGTGCGCCGCGGCAAGCCGCGGCGGCACTAGGTCA 49						
Db	52 CATTCAACCGAGAGCCGCCCGAGCAATCCAGAGTCCCGCTCGCTCA 7						
RESULT 8	1115 bp						
LOCUS	BM5474535/c						
DEFINITION	602517304F1 NIH_MGC_16 Homo sapiens cDNA clone IMAGE:4649112 5',						
ACCESSION	BM5474535						
VERSION	BM5474535.1	GI:13406812					
KEYWORDS	EST.						
SOURCE	human.						
ORGANISM	Homo sapiens						
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;						
	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.						
TITLE	NIH-MGC http://mhc.nci.nih.gov/.						
AUTHORS	1 (bases 1 to 1208)						
JOURNAL	National Institutes of Health, Mammalian Gene Collection (MGC)						
COMMENT	Unpublished (1999)						
	Contact: Robert Strausberg, Ph.D.						

Email: c9apbs-remail.nlh.gov
Tissue Procurement: AFCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
plate: LLCM1428 row: g column: 01
High quality sequence stop: 478.

FEATURES
SOURCE
1. 1208
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:4649112"
/clone_lib="NIH_MGC_16"
/tissue_type="retinoblastoma"
/lab_host="DH10B (phage-resistant)"
/note="Organ: eye; Vector: pORF7; Site_1: XhoI; Site_2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH_MGC library."

BASE COUNT
ORIGIN
361 a 349 c 313 g 185 t

Query Match 42.9% Score 23.6 DB 12 Length 1208;
Best Local Similarity 69.6% Pred No. 7e+02; Indels 0; Gaps 0;
Matches 32; Conservative 0; Mismatches 14;

QY 1 GTCATTGACCTAGTCCCGCCGCAAGCCCGGCGGCATAGG 46
||||||| | | | | | | | | | | | | | | | | |
DB 1008 GTACATTGCCCTGTCTCCCGCGCTCAGCACGCGTGCACGTGG 963

RESULT 9 596 bp mRNA linear EST 21-APR-2001
Bg194272
LOCUS Bg194272
DEFINITION RST13417 Atherys RAGE Library Homo sapiens CDNA, mRNA sequence.
ACCESSION Bg194272
VERSION Bg194272.1 GI:13715959
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 596)
Harrington,J.J., Sherf,B., Rundlett,S., Jackson,P.D., Perry,R.,
Cain,S., Leventhal,C., Thonon,M., Ramachandran,R., Whittington,J.
Lerner,L., Costanzo,D., McEllisolt,K., Boxer,S., Mays,R., Smith
E., Veloso,N., Kliska,A., Hess,J., Colhren,K., Lo,K., Offenbacher
J., Danzig,J. and Ducar,M.
Creation of genome-wide protein expression libraries using random
activation of gene expression
Nat. Biotechnol. 19 (5), 440-445 (2001)
21227151
Contact: Scott J. Cain
Atherys, Inc.
3201 Carnegie Ave. Cleveland, OH 44115, USA
Tel: 216 431 9900
Fax: 216 361 9596
Email: scaine@atherys.com
High quality sequence stop: 596.
Location/Qualifiers
1. 596
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Atherys RAGE Library"
/cell_line="HT1080"
/note="See 'Creation of Genome-wide Protein Expression
Libraries using Random Activation of Gene Expression',

FEATURES
SOURCE

Nature Biotechnology, in press. Note that even though the cell type indicated is HT1080, since a random activation method was used, these sequence tags are not necessarily expressed in HT1080 under normal circumstances."

BASE COUNT 181 a 140 c 126 g 148 t 1 others

ORIGIN

Query Match 42.5% Score 23.4; DB 12; Length 596;
Best Local Similarity 67.3% Pred. No. 6.8e+02;
Matches 33; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

0Y 4 CATTGACCTAGTGGCCGCGGCAAGCCGCGGACATGCAATC 52
1 ||||| 11 111 111 111 111 111 111 111 111 111
Db 6 CATTGACGAGGTACCTGGCGACGACGAGGACCACTAGCCCAAC 54

RESULT 10
BO892955

BO892955 884 bp mRNA linear EST 16-AUG-2002
AGENCOURT 8124301 lupskl_dorsal_root_ganglion Homo sapiens cDNA
clone IMAGE:6178096 5', mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BO892955
BO892955
BO892955.1 GI:22284965
EST.
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

NIH-MGC http://mgc.nci.nih.gov/
1 (bases 1 to 884)
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov

FEATURES
source

Tissue Procurement: Dr. James R. Lupski
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
http://image.llnl.gov
Plate: LLM13557 row: b column: 17
High quality sequence stop: 629.

FEATURES
source

1. 884
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:6178096"
/clone_11b="lupskl_dorsal_root_ganglion"
/sex="male"
/tissue_type="dorsal root ganglia"
/dev_stage="adult" 36 yr
/lab_host="DH10B"
/note="Vector: pCMV-SPORT6 (Life Technologies); Site_1:
NotI; Site_2: SalI; cDNA made by oligo-dT priming.
Directionally cloned using the following adaptors:
5'-TCGACCCACGCGTCCG-3' and
5'-GACCTAGTCTAGATGCGGAGCGGCGGCGGCT(15)-3'. Size selected >
1 kb for average insert length 1.7 kb. This is a primary
library, non-amplified. Library constructed by Life
Technologies and donated by J. Lupski, M.D./Ph.D. (Baylor
College of Medicine) and is available through Life
Technologies."

BASE COUNT 226 a 215 c 335 g 108 t

ORIGIN

Query Match 42.5% Score 23.4; DB 14; Length 884;
Best Local Similarity 73.2% Pred. No. 7.5e+02;
Matches 30; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

0Y 1 GTACATGACCTAGTGGCCGCGGCAAGCCGCGGACGCA 41
1 ||||| 111 111 111 111 111 111 111 111 111 111
Db 826 GGACATGCGCCTGGACCCCGCGGCAATCCGGGGGCA 866

BASE COUNT 226 a 215 c 335 g 108 t

ORIGIN

Query Match 42.5% Score 23.4; DB 14; Length 884;
Best Local Similarity 73.2% Pred. No. 7.5e+02;
Matches 30; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

0Y 1 GTACATGACCTAGTGGCCGCGGCAAGCCGCGGACGCA 41
1 ||||| 111 111 111 111 111 111 111 111 111 111
Db 826 GGACATGCGCCTGGACCCCGCGGCAATCCGGGGGCA 866

BASE COUNT 226 a 215 c 335 g 108 t

ORIGIN

Query Match 42.5% Score 23.4; DB 14; Length 884;
Best Local Similarity 73.2% Pred. No. 7.5e+02;
Matches 30; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

0Y 1 GTACATGACCTAGTGGCCGCGGCAAGCCGCGGACGCA 41
1 ||||| 111 111 111 111 111 111 111 111 111 111
Db 826 GGACATGCGCCTGGACCCCGCGGCAATCCGGGGGCA 866

BASE COUNT 226 a 215 c 335 g 108 t

ORIGIN

RESULT 11
BO642928

BO642928 1149 bp mRNA linear EST 15-JUL-2002
AGENCOURT_8485188 NIH-MGC_99 Homo sapiens cDNA clone IMAGE:6305160
5', mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BO642928
BO642928.1 GI:21767100
EST.
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

NIH-MGC http://mgc.nci.nih.gov/
1 (bases 1 to 1149)
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov

FEATURES
source

Tissue Procurement: Lou Staudt
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
http://image.llnl.gov
Plate: LLM2527 row: a column: 01
High quality sequence start: 33
High quality sequence stop: 327.

FEATURES
source

1. 1149
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:6305160"
/clone_11b="NIH-MGC_99"
/tissue_type="lymphoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: lymph; Vector: pOTB7; Site_1: XhoI; Site_2:
EcoRI; cDNA made by oligo-dT priming. Directionally cloned
into EcoRI/XhoI sites using the following 5' adaptor:
GGCAGCAG(G). Size-selected >500bp for average insert size
1.8kb. Library constructed by Ling Hong in the laboratory
of Gerald M. Rubin (University of California, Berkeley)
using ZAP-cDNA synthesis kit (Stratagene) and Superscript
II RT (Life Technologies). Note: this is a NIH-MGC
Library."

BASE COUNT 162 a 442 c 398 g 140 t 7 others

ORIGIN

Query Match 42.5% Score 23.4; DB 14; Length 1149;
Best Local Similarity 73.2% Pred. No. 8e+02;
Matches 30; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

0Y 3 ACATGACCTAGTGGCCGCGGCAAGCCGCGGACGCACT 43
1 ||||| 111 111 111 111 111 111 111 111 111 111
Db 453 ACTTGGCGGAGTGGCCGCGGCAAGCCGCGGCGGCGGCT 493

BASE COUNT 162 a 442 c 398 g 140 t 7 others

ORIGIN

Query Match 42.5% Score 23.4; DB 14; Length 1149;
Best Local Similarity 73.2% Pred. No. 8e+02;
Matches 30; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

0Y 3 ACATGACCTAGTGGCCGCGGCAAGCCGCGGACGCACT 43
1 ||||| 111 111 111 111 111 111 111 111 111 111
Db 453 ACTTGGCGGAGTGGCCGCGGCAAGCCGCGGCGGCGGCT 493

BASE COUNT 162 a 442 c 398 g 140 t 7 others

ORIGIN

Query Match 42.5% Score 23.4; DB 14; Length 1149;
Best Local Similarity 73.2% Pred. No. 8e+02;
Matches 30; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

0Y 3 ACATGACCTAGTGGCCGCGGCAAGCCGCGGACGCACT 43
1 ||||| 111 111 111 111 111 111 111 111 111 111
Db 453 ACTTGGCGGAGTGGCCGCGGCAAGCCGCGGCGGCGGCT 493

BASE COUNT 162 a 442 c 398 g 140 t 7 others

ORIGIN

Query Match 42.5% Score 23.4; DB 14; Length 1149;
Best Local Similarity 73.2% Pred. No. 8e+02;
Matches 30; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

0Y 3 ACATGACCTAGTGGCCGCGGCAAGCCGCGGACGCACT 43
1 ||||| 111 111 111 111 111 111 111 111 111 111
Db 453 ACTTGGCGGAGTGGCCGCGGCAAGCCGCGGCGGCGGCT 493

BASE COUNT 162 a 442 c 398 g 140 t 7 others

ORIGIN

DIVISION OF BIOLOGY 150-25
California Institute of Technology
Pasadena California 91125, USA

Cap-trapper method (Caininci, in preparation). Library constructed by M. Brownstein (NIMH/NIHGR1, National Institutes of Health). Note: this is a NIH-MGC Library.

BASE COUNT 133 a 193 c 148 g 159 t
ORIGIN

Query Match 42.2%; Score 23.2; DB 13; Length 633;
Best Local Similarity 70.5%; Pred. No. 8e+02;
Matches 31; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 3 ACATTGACTACTGCGCGCGCAAGCCGCGCGGCGACTAGG 46
DB 30 ACAGTCCCGCAGCGCGCGCGCGCGCGCGCGCTCTAGG 73

RESULT 15
B1669165 682 bp mRNA linear EST 12-SEP-2001
LOCUS 603293596F1 NIH_MGC_96 Homo sapiens CDNA clone IMAGE:5314822 5',
DEFINITION mRNA sequence.

ACCESSION B1669165
VERSION B1669165
KEYWORDS B1669165.1 GI:15583398
SOURCE EST.
ORGANISM human.

REFERENCE Homo sapiens
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
JOURNAL NIH-MGC http://mgc.nci.nih.gov/
COMMENT Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Miklos Palokovits, M.D., Ph.D.
CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
Toshiyuki and Piero Carninci (RIKEN)
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LMNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/ILNL at:
http://image.llnl.gov
Plate: L1AM11797 row: P. Column: 23
High quality sequence stop: 682.
Location/Qualifiers

FEATURES
SOURCE 1..682
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5314822"
/clone_lib="NIH_MGC_96"
/tissue_type="hypothalamus"
/lab_host="DH10B"
/note="Organ: brain; Vector: pBluescriptR (modified
pBluescript KS+); Site_1: BamHI; Site_2: SalI-xhoI (gtcgag
); Oligo-dT primed using primer 5'-TTTTTTTTTTTTTTVN-3',
size-selected for average insert size 2.3 kb and
normalized to 10^5. This is a primary library enriched
for full-length clones and constructed using the
Cap-trapper method (Carninci, in preparation). Library
constructed by M. Brownstein (NIH/NHGRI, National
Institutes of Health). Note: this is a NIH_MGC Library."

BASE COUNT 159 a 176 c 184 g 163 t
ORIGIN

Query Match 42.2%; Score 23.2; DB 13; Length 682;
Best Local Similarity 70.5%; Pred. No. 8.2e+02;
Matches 31; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 3 ACATTGACTACTGCGCGCGCAAGCCGCGCGGCGACTAGG 46
DB 73 ACAGTCCCGCAGCGCGCGCGCGCGCGCGCTCTAGG 116

Search completed: December 27, 2002, 06:15:34
Job time : 1327.5 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 27, 2002, 03:13:39 : Search time 1330 Seconds

(without alignments)
1203.499 Million cell updates/sec

Title: US-09-918-029-20

Perfect score: 55
Sequence: 1 gtcacatcgcttgacctaagt.....ccggcgcgacctaagtcaat 55

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : GenBank:*

1: gb_da:*
2: gb_htg:*
3: gb_in:*
4: gb_cm:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vl:*
15: em_da:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_cm:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vl:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htg_mus:*
34: em_htg_pln:*
35: em_htg_rod:*
36: em_htg_mam:*
37: em_htg_vrt:*
38: em_sy:*
39: em_htgo_hum:*
40: em_htgo_mus:*
41: em_htgo_other:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	55	100.0	55	6 AR091517	AR091517 Sequence
2	55	100.0	55	6 AR154397	AR154397 Sequence
3	55	100.0	55	6 AR174325	AR174325 Sequence
4	55	100.0	55	6 AR183309	AR183309 Sequence
5	55	100.0	55	6 AR080361	AR080361 Sequence
6	51	92.7	55	6 AR091516	AR091516 Sequence
7	51	92.7	55	6 AR154396	AR154396 Sequence
8	51	92.7	55	6 AR174324	AR174324 Sequence
9	51	92.7	55	6 AR183308	AR183308 Sequence
10	51	92.7	55	6 AX080360	AX080360 Sequence
11	43.4	78.9	50	6 AR091514	AR091514 Sequence
12	43.4	78.9	50	6 AR091515	AR091515 Sequence
13	43.4	78.9	50	6 AR154394	AR154394 Sequence
14	43.4	78.9	50	6 AR154395	AR154395 Sequence
15	43.4	78.9	50	6 AR174322	AR174322 Sequence
16	43.4	78.9	50	6 AR174323	AR174323 Sequence
17	43.4	78.9	50	6 AR183306	AR183306 Sequence
18	43.4	78.9	50	6 AR183307	AR183307 Sequence
19	43.4	78.9	50	6 AX080358	AX080358 Sequence
20	43.4	78.9	50	6 AX080359	AX080359 Sequence
21	42.2	76.7	55	6 AR091516	AR091516 Sequence
22	42.2	76.7	55	6 AR091517	AR091517 Sequence
23	42.2	76.7	55	6 AR154396	AR154396 Sequence
24	42.2	76.7	55	6 AR154397	AR154397 Sequence
25	42.2	76.7	55	6 AR174324	AR174324 Sequence
26	42.2	76.7	55	6 AR174325	AR174325 Sequence
27	42.2	76.7	55	6 AR183308	AR183308 Sequence
28	42.2	76.7	55	6 AR183309	AR183309 Sequence
29	42.2	76.7	55	6 AX080360	AX080360 Sequence
30	42.2	76.7	55	6 AX080361	AX080361 Sequence
31	39.4	71.6	45	6 AR091519	AR091519 Sequence
32	39.4	71.6	45	6 AR154399	AR154399 Sequence
33	39.4	71.6	45	6 AR174327	AR174327 Sequence
34	39.4	71.6	45	6 AR183314	AR183314 Sequence
35	38.6	70.2	50	6 AR091514	AR091514 Sequence
36	38.6	70.2	50	6 AR091515	AR091515 Sequence
37	38.6	70.2	50	6 AR154394	AR154394 Sequence
38	38.6	70.2	50	6 AR154395	AR154395 Sequence
39	38.6	70.2	50	6 AR174322	AR174322 Sequence
40	38.6	70.2	50	6 AR174323	AR174323 Sequence
41	38.6	70.2	50	6 AR183306	AR183306 Sequence
42	38.6	70.2	50	6 AR183307	AR183307 Sequence
43	38.6	70.2	50	6 AX080358	AX080358 Sequence
44	38.6	70.2	50	6 AX080359	AX080359 Sequence
45	36.2	65.8	45	6 AX080363	AX080363 Sequence

ALIGNMENTS

RESULT 1
LOCUS AR091517 55 bp DNA linear PAT 07-SEP-2000
DEFINITION Sequence 20 from patent US 5994128.
ACCESSION AR091517
VERSION AR091517.1 GI:10018272
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE Unclassified.
1 (bases 1 to 55)
AUTHORS Fallaux,F.Jacobsus., Hoebein,R.Cornellis., Van der Eb,A.Jan., Boul,A.
and Valerio,D.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy

JOURNAL Patent: US 5994128-A 20 30-NOV-1999;
FEATURES Location/Qualifiers
source 1..55
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN

Query Match 100.0%; Score 55; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 2.4e-08;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55
Db 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55

RESULT 2
LOCUS AR154397 55 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 20 from patent US 6238893.
ACCESSION AR154397
VERSION AR154397.1 GI:15122450
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 55)
AUTHORS Hoeber, R. Cornelis, and Boul, A.
TITLE Method for intracellular DNA amplification
JOURNAL Patent: US 6238893-A 20 29-MAY-2001;
FEATURES Location/Qualifiers
source 1..55
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN

Query Match 100.0%; Score 55; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 2.4e-08;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55
Db 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55

RESULT 3
LOCUS AR174325 55 bp DNA linear PAT 17-DEC-2001
DEFINITION Sequence 20 from patent US 6306652.
ACCESSION AR174325
VERSION AR174325.1 GI:17914645
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 55)
AUTHORS Fallaux, F. Jacobs, Hoeber, R. Cornelis, Van Der Eb, A. Jan., Boul, A. and Valerio, D.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL Patent: US 6306652-A 20 23-OCT-2001;
FEATURES Location/Qualifiers
source 1..55
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN

Query Match 100.0%; Score 55; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 2.4e-08;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55
Db 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55

Db 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55

RESULT 4
LOCUS AR183309 55 bp DNA linear PAT 20-APR-2002
DEFINITION Sequence 42 from patent US 6340595.
ACCESSION AR183309
VERSION AR183309.1 GI:20226902
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 55)
AUTHORS Vogels, R., Boul, A., van Es, H. and Schouten, G.
TITLE High throughput screening of gene function using adenoviral libraries for functional genomics applications
JOURNAL Patent: US 6340595-A 42 22-JAN-2002;
FEATURES Location/Qualifiers
source 1..55
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN

Query Match 100.0%; Score 55; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 2.4e-08;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55
Db 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55

RESULT 5
LOCUS AX080361 55 bp DNA linear PAT 22-FEB-2001
DEFINITION Sequence 20 from Patent WO0105945.
ACCESSION AX080361
VERSION AX080361.1 GI:13159819
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 55)
AUTHORS Hoeber, R. C., Boul, A., Valerio, D., van der Eb, A. J., Schouten, G. and Fallaux, F. J.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL Patent: WO 0105945-A 20 25-JAN-2001;
FEATURES Location/Qualifiers
source 1..55
BASE COUNT 9 a 17 c 17 g 12 t
ORIGIN

Query Match 100.0%; Score 55; DB 6; Length 55;
Best Local Similarity 100.0%; Pred. No. 2.4e-08;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55
Db 1 GTACATCGATTGACCTAGTGCCTGGGCTTTGGCCGGGGGCGGCACTAGTCAAT 55

RESULT 6
LOCUS AR091516/c 55 bp DNA linear PAT 07-SEP-2000
DEFINITION Sequence 19 from patent US 5994128.
ACCESSION AR091516
VERSION AR091516.1 GI:10018271

KEYWORDS	Unknown.
SOURCE	Unknown.
ORGANISM	Unclassified.
REFERENCE	1 (bases 1 to 55)
AUTHORS	Fallaux,F.Jacobus., Hoeben,R.Cornelis., Van der Eb,A.Jan., Bout,A. and Valerio,D.
TITLE	Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL	Patent: US 5994128-A 19 30-NOV-1999;
FEATURES	Location/Qualifiers
source	1..55
BASE COUNT	12 a 17 c 17 g 9 t
ORIGIN	/organism="unknown"
Query Match	92.7%; Score 51; DB 6; Length 55;
Best Local Similarity	100.0%; Pred.No. 5.5e-07;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
OY	5 ATCGATTGACCTAGTGCAGCGCGGCTTTGCCCGGCGGCACTAGTCAAT 55
Db	55 ATCGATTGACCTAGTGCAGCGCGGCTTTGCCCGGCGGCACTAGTCAAT 5
RESULT 7	
LOCUS	ARI54396/c 55 bp DNA linear PAT 08-AUG-2001
DEFINITION	Sequence 19 from patent US 6238893.
ACCESSION	ARI54396
VERSION	ARI54396.1 GI:15122449
KEYWORDS	.
SOURCE	Unknown.
ORGANISM	Unclassified.
REFERENCE	1 (bases 1 to 55)
AUTHORS	Hoeben,R.Cornelis, and Bout,A.
TITLE	Method for Intracellular DNA amplification
JOURNAL	Patent: US 6238893-A 19 29-MAY-2001;
FEATURES	Location/Qualifiers
source	1..55
BASE COUNT	12 a 17 c 17 g 9 t
ORIGIN	/organism="unknown"
Query Match	92.7%; Score 51; DB 6; Length 55;
Best Local Similarity	100.0%; Pred.No. 5.5e-07;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
OY	5 ATCGATTGACCTAGTGCAGCGCGGCTTTGCCCGGCGGCACTAGTCAAT 55
Db	55 ATCGATTGACCTAGTGCAGCGCGGCTTTGCCCGGCGGCACTAGTCAAT 5
RESULT 8	
LOCUS	ARI74324/c 55 bp DNA linear PAT 17-DEC-2001
DEFINITION	Sequence 19 from patent US 6306652.
ACCESSION	ARI74324
VERSION	ARI74324.1 GI:17914644
KEYWORDS	.
SOURCE	Unknown.
ORGANISM	Unclassified.
REFERENCE	1 (bases 1 to 55)
AUTHORS	Fallaux,F.Jacobus., Hoeben,R.Cornelis., Van Der Eb,A.Jan., Bout,A. and Valerio,D.
TITLE	Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL	Patent: US 6306652-A 19 23-OCT-2001;
FEATURES	Location/Qualifiers
source	1..55
	/organism="unknown"

ORIGIN	BASE COUNT	12 a	17 c	17 g	9 t
Query Match	92.7%;	Score 51;	DB 6;	Length 55;	
Best Local Similarity	100.0%;	Pred. No. 5.5e-07;			
Matches	51;	Conservative	0;	Mismatches	0;
				Indels	0;
				Gaps	0;
Oy	5	ATCGATTGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACTAGTCAAT	55		
Db	55	ATCGATTGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACTAGTCAAT	5		
RESULT 9					
LOCUS	ARI83308/c		55 bp	DNA	Linear
DEFINITION	Sequence	41	from patent US 6340595.		PAT 20-APR-2002
ACCESSION	ARI83308				
VERSION	ARI83308.1		GI:20226901		
KEYWORDS					
SOURCE	Unknown.				
ORGANISM	Unknown.				
REFERENCE	1 (bases 1 to 55)				
AUTHORS	Vogels,R., Bout,A., van Es,H. and Schouten,G.				
TITLE	High throughput screening of gene function using adenoviral				
JOURNAL	libraries for functional genomics applications				
FEATURES	Patent: US 6340595-A 41 22-JAN-2002;				
	Location/Qualifiers				
source	1..55				
BASE COUNT	12 a	17 c	17 g	9 t	
ORIGIN	/organism="Unknown"				
Query Match	92.7%;	Score 51;	DB 6;	Length 55;	
Best Local Similarity	100.0%;	Pred. No. 5.5e-07;			
Matches	51;	Conservative	0;	Mismatches	0;
				Indels	0;
				Gaps	0;
Oy	5	ATCGATTGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACTAGTCAAT	55		
Db	55	ATCGATTGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACTAGTCAAT	5		
RESULT 10					
LOCUS	AX080360/c		55 bp	DNA	Linear
DEFINITION	Sequence	19	from Patent W00105945.		PAT 22-FEB-2001
ACCESSION	AX080360				
VERSION	AX080360.1		GI:13159818		
KEYWORDS					
SOURCE	synthetic construct.				
ORGANISM	synthetic construct.				
REFERENCE	artificial sequences.				
AUTHORS	1 (bases 1 to 55)				
	Hoeben,R.C., Boul,A., Valerio,D., van der Eb,A.J., Schouten,G. and				
	Fallaux,F.J.				
TITLE	Packaging systems for human recombinant adenovirus to be used in				
JOURNAL	gene therapy				
	patent: WO 0105945-A 19 25-JAN-2001;				
	Introgene B.V. (NL)				
FEATURES	Location/Qualifiers				
source	1..55				
BASE COUNT	12 a	17 c	17 g	9 t	
ORIGIN	/organism="synthetic construct"				
	/db xref="taxon:32630"				
	/note="oligonucleotide HP/cial"				
Query Match	92.7%;	Score 51;	DB 6;	Length 55;	
Best Local Similarity	100.0%;	Pred. No. 5.5e-07;			
Matches	51;	Conservative	0;	Mismatches	0;
				Indels	0;
				Gaps	0;
Oy	5	ATCGATTGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACTAGTCAAT	55		

Db 55 ATGCATTGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 5

RESULT 11
LOCUS AR091514/c 50 bp DNA linear PAT 07-SEP-2000
DEFINITION Sequence 17 from patent US 5994128.
ACCESSION AR091514
VERSION AR091514.1 GI:10018269
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 50)
AUTHORS Fallaux,F.,Jacobus., Hoeben,R.,Cornelis., Van der Eb,A.,Jan., Bout,A.
and Valerio,D.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL Patent: US 5994128-A 17 30-NOV-1999;
FEATURES
source location/Qualifiers
1..50
/organism="unknown"
BASE COUNT 10 a 17 c 17 g 6 t

Query Match 78.9%; Score 43.4; DB 6; Length 50;
Best Local Similarity 97.8%; Pred. No. 0.00021;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 11 TGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 55
Db 49 TGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 5

RESULT 12
LOCUS AR091515 50 bp DNA linear PAT 07-SEP-2000
DEFINITION Sequence 18 from patent US 5994128.
ACCESSION AR091515
VERSION AR091515.1 GI:10018270
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 50)
AUTHORS Fallaux,F.,Jacobus., Hoeben,R.,Cornelis., Van der Eb,A.,Jan., Bout,A.
and Valerio,D.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL Patent: US 5994128-A 18 30-NOV-1999;
FEATURES
source location/Qualifiers
1..50
/organism="unknown"
BASE COUNT 6 a 17 c 17 g 10 t

Query Match 78.9%; Score 43.4; DB 6; Length 50;
Best Local Similarity 97.8%; Pred. No. 0.00021;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 11 TGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 55
Db 6 TGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 50

RESULT 13
LOCUS AR154394/c 50 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 17 from patent US 6238893.
ACCESSION AR154394
VERSION AR154394.1 GI:15122447
KEYWORDS
SOURCE Unknown.

ORGANISM Unknown.
REFERENCE 1 (bases 1 to 50)
AUTHORS Hoeben,R.,Cornelis., and Bout,A.
TITLE Method for intracellular DNA amplification
JOURNAL Patent: US 6238893-A 17 29-MAY-2001;
FEATURES
source location/Qualifiers
1..50
/organism="unknown"
BASE COUNT 10 a 17 c 17 g 6 t

Query Match 78.9%; Score 43.4; DB 6; Length 50;
Best Local Similarity 97.8%; Pred. No. 0.00021;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 11 TGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 55
Db 49 TGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 5

RESULT 14
LOCUS AR154395 50 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 18 from patent US 6238893.
ACCESSION AR154395
VERSION AR154395.1 GI:15122448
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 50)
AUTHORS Hoeben,R.,Cornelis., and Bout,A.
TITLE Method for intracellular DNA amplification
JOURNAL Patent: US 6238893-A 18 29-MAY-2001;
FEATURES
source location/Qualifiers
1..50
/organism="unknown"
BASE COUNT 6 a 17 c 17 g 10 t

Query Match 78.9%; Score 43.4; DB 6; Length 50;
Best Local Similarity 97.8%; Pred. No. 0.00021;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 11 TGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 55
Db 6 TGACCTAGTCCGCCGGCGCTTTCGCCCGCGGCACACTAGTCAAT 50

RESULT 15
LOCUS AR174322/c 50 bp DNA linear PAT 17-DEC-2001
DEFINITION Sequence 17 from patent US 630652.
ACCESSION AR174322
VERSION AR174322.1 GI:17914642
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 50)
AUTHORS Fallaux,F.,Jacobus., Hoeben,R.,Cornelis., Van Der Eb,A.,Jan., Bout,A.
and Valerio,D.
TITLE Packaging systems for human recombinant adenovirus to be used in gene therapy
JOURNAL Patent: US 630652-A 17 23-OCT-2001;
FEATURES
source location/Qualifiers
1..50
/organism="unknown"
BASE COUNT 10 a 17 c 17 g 6 t

Query Match 78.9%; Score 43.4; DB 6; Length 50;

Best Local Similarity 97.8%; Pred. No. 0.00021;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 11 TGACCTAGTGGCCGCCGGGCTTTGGCCGGGGGCACTAGTCAAT 55
|||||
Db 49 TGACCTAGTGGCCGCCGGGCTTTGGCCGGGGGCACTAGTCAAGT 5

Search completed: December 27, 2002, 05:31:03
Job time : 1332.5 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 27, 2002, 04:38:45 ; Search time 33 Seconds
(without alignments)
511.128 Million cell updates/sec

Title: US-09-918-029-20

Perfect score: 55
Sequence: 1 gtaacatgattgactagtg.....ccgggagcgactagtcacat 55

Scoring table: IDENTITY_MTC
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 15338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued_Patents_NA:*
1: /cgn2_6/prodata/1/lna/5A_COMB.seq:*
2: /cgn2_6/prodata/1/lna/5B_COMB.seq:*
3: /cgn2_6/prodata/1/lna/5A_COMB.seq:*
4: /cgn2_6/prodata/1/lna/5B_COMB.seq:*
5: /cgn2_6/prodata/1/lna/PCUTUS_COMB.seq:*
6: /cgn2_6/prodata/1/lna/backfillseq1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query	Match length	DB	ID	Description
No.	Score				
1	55	100.0	55	2	US-08-793-170-20
2	55	100.0	55	3	US-08-892-873-20
3	55	100.0	55	4	US-09-334-765A-20
4	55	100.0	55	4	US-09-356-575E-20
5	55	100.0	55	4	US-09-333-820-20
6	55	100.0	55	4	US-09-358-036-42
7	55	100.0	55	4	US-09-097-239-42
8	55	100.0	55	2	US-08-793-170-19
9	51	92.7	55	3	US-08-892-873-19
10	51	92.7	55	4	US-09-334-765A-19
11	51	92.7	55	4	US-09-356-575E-19
12	51	92.7	55	4	US-09-333-820-19
13	51	92.7	55	4	US-09-358-036-41
14	51	92.7	55	4	US-09-097-239-41
15	43.4	78.9	50	2	US-08-793-170-17
16	43.4	78.9	50	2	US-08-793-170-18
17	43.4	78.9	50	3	US-08-892-873-17
18	43.4	78.9	50	3	US-08-892-873-18
19	43.4	78.9	50	4	US-09-334-765A-17
20	43.4	78.9	50	4	US-09-334-765A-18
21	43.4	78.9	50	4	US-09-356-575E-17
22	43.4	78.9	50	4	US-09-356-575E-18
23	43.4	78.9	50	4	US-09-333-820-17
24	43.4	78.9	50	4	US-09-333-820-18
25	43.4	78.9	50	4	US-09-358-036-39
26	43.4	78.9	50	4	US-09-358-036-40
27	43.4	78.9	50	4	US-09-097-239-39

28	43.4	78.9	50	4	US-09-097-239-40	Sequence 40, Appl
29	42.2	76.7	55	2	US-08-793-170-19	Sequence 19, Appl
30	42.2	76.7	55	2	US-08-793-170-20	Sequence 20, Appl
31	42.2	76.7	55	3	US-08-892-873-19	Sequence 19, Appl
32	42.2	76.7	55	3	US-08-892-873-20	Sequence 20, Appl
33	42.2	76.7	55	4	US-09-334-765A-19	Sequence 19, Appl
34	42.2	76.7	55	4	US-09-334-765A-20	Sequence 20, Appl
35	42.2	76.7	55	4	US-09-356-575E-19	Sequence 19, Appl
36	42.2	76.7	55	4	US-09-356-575E-20	Sequence 20, Appl
37	42.2	76.7	55	4	US-09-333-820-19	Sequence 19, Appl
38	42.2	76.7	55	4	US-09-333-820-20	Sequence 20, Appl
39	42.2	76.7	55	4	US-09-358-036-41	Sequence 41, Appl
40	42.2	76.7	55	4	US-09-358-036-42	Sequence 42, Appl
41	42.2	76.7	55	4	US-09-097-239-41	Sequence 41, Appl
42	42.2	76.7	55	4	US-09-097-239-42	Sequence 42, Appl
43	39.4	71.6	45	3	US-08-793-170-22	Sequence 22, Appl
44	39.4	71.6	45	3	US-08-892-873-22	Sequence 22, Appl
45	39.4	71.6	45	4	US-09-334-765A-22	Sequence 22, Appl

ALIGNMENTS

RESULT 1
US-08-793-170-20
Sequence 20, Application US/08793170
Patent No. 5994128
GENERAL INFORMATION:
APPLICANT: FALLAUX et al.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
NUMBER OF INVENTION: ADENOVIRUS TO BE USED IN GENE THERAPY
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESSES:
ADDRESSEE: RAE-VENTER LAW GROUP, P.C.
STREET: 260 SHERIDAN AVENUE, PO BOX 60039
CITY: PALO ALTO
STATE: CALIFORNIA
COUNTRY: USA
ZIP: 94306
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/793,170
FILING DATE: 25-MAR-1997
CLASSIFICATION: 435
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: WO 97/00326
FILING DATE: 14-JUN-1996
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: EP 95201728.3
FILING DATE: 26-JUN-1995
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: EP 95201611.1
FILING DATE: 15-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: RAE-VENTER, BARBARA
REGISTRATION NUMBER: 32,750
REFERENCE/DOCKET NUMBER: INGE.002.00US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650)328-4400
TELEFAX: (650)328-3377
TELEX: N/A
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 55 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
US-08-793-170-20


```

: FEATURE:
: OTHER INFORMATION: Derived from Adenovirus
US-09-356-575E-20

Query Match          100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 5,7e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GTACATCGATTGACCTAGTCCGCCCGGGCTTTGCCCGGGCGGACACTAGTCAAT 55
        |||||||
Db      1 GTACATCGATTGACCTAGTCCGCCCGGGCTTTGCCCGGGCGGACACTAGTCAAT 55

RESULT 5
US-09-333-820-20
: Sequence 20, Application US/09333820A
: Patent No. 6306552
: GENERAL INFORMATION:
: APPLICANT: Fallaux, Frits J.
: APPLICANT: Hoeber, Robert C.
: APPLICANT: Boul, Abraham
: APPLICANT: Valerio, Domenico
: APPLICANT: Van der Eb, Alex J.
: TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO BE USED IN
: FILE REFERENCE: 3833.1US
: CURRENT APPLICATION NUMBER: US/09/333,820A
: EARLIER FILING DATE: 1999-06-15
: EARLIER APPLICATION NUMBER: US 08/793,170
: EARLIER FILING DATE: 1997-03-25
: EARLIER APPLICATION NUMBER: PCT/NL96/00244
: EARLIER FILING DATE: 1996-06-14
: EARLIER APPLICATION NUMBER: EP 95201728.3
: EARLIER FILING DATE: 1995-06-26
: EARLIER APPLICATION NUMBER: EP 95201611.1
: EARLIER FILING DATE: 1995-06-15
: NUMBER OF SEQ ID NOS: 22
: SOFTWARE: Corel WordPerfect 8.0
: SEQ ID NO 20
: LENGTH: 55
: TYPE: DNA
: ORGANISM: Artificial Sequence
: FEATURE:
: NAME/KEY:
: LOCATION:
: OTHER INFORMATION: Description of Artificial Sequence: primer HP/c1a2
US-09-333-820-20

Query Match          100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 5,7e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GTACATCGATTGACCTAGTCCGCCCGGGCTTTGCCCGGGCGGACACTAGTCAAT 55
        |||||||
Db      1 GTACATCGATTGACCTAGTCCGCCCGGGCTTTGCCCGGGCGGACACTAGTCAAT 55

RESULT 6
US-09-358-036-42
: Sequence 42, Application US/09358036
: Patent No. 6340595
: GENERAL INFORMATION:
: APPLICANT: Vogels, Ronald
: APPLICANT: Boul, Abraham
: APPLICANT: van Es, Helmut
: APPLICANT: Schouten, Govert
: TITLE OF INVENTION: High Throughput Screening of Gene Function Using
: TITLE OF INVENTION: Adenoviral Libraries for Functional Genomics
: FILE REFERENCE: 21834108
: CURRENT APPLICATION NUMBER: US/09/358,036
: EARLIER FILING DATE: 1999-07-21
: EARLIER APPLICATION NUMBER: US 09/097,239

```

```

      EARLIER FILING DATE: 1995-07-25
      NUMBER OF SEQ ID NOS: 69
      SOFTWARE: PatentIn Ver. 2.0
      SEQ ID NO 42
      LENGTH: 55
      TYPE: DNA
      ORGANISM: Artificial Sequence
      FEATURE:
      OTHER INFORMATION: Description of Artificial Sequence:oligonucleotide
US-09-358-036-42

Query Match                      100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 5,7e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      1  GTCATCGATTGACCTAGTGCCTGGCCGGGCTTTGGCCGGCGGACATAGAGTCAT 55
        |||||||
DB      1  GTCATCGATTGACCTAGTGCCTGGCCGGGCTTTGGCCGGCGGACATAGAGTCAT 55

RESULT 7
US-09-097-239-42
; Sequence 42, Application US/09097239
; Patent No. 6413776
; GENERAL INFORMATION:
; APPLICANT: VOGELS, RONALD,
; APPLICANT: BOOT, ABRAHAM,
; APPLICANT: VAN ES, HELMUTH HG,
; APPLICANT: SCHOUTEN, GOVERT
; TITLE OF INVENTION: HIGH THROUGHPUT SCREENING OF GENE
; TITLE OF INVENTION: FUNCTION USING ADENOVIARAL LIBRARIES FOR FUNCTIONAL
; NUMBER OF SEQUENCES: 47
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: RAE-VENTER LAW GROUP
; STREET: PO BOX 60039
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94306
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/097,239
; FILING DATE: 12-JUN-1998
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: RAE-VENTER, BARBARA
; REGISTRATION NUMBER: 32,750
; REFERENCE/DOCKET NUMBER: INGE.008.000S
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650)328-4400
; TELEFAX: (650)328-4477
; INFORMATION FOR SEQ ID NO: 42:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 55 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
;
;
US-09-097-239-42

Query Match                      100.0%; Score 55; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 5,7e-11;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      1  GTCATCGATTGACCTAGTGCCTGGCCGGGCTTTGGCCGGCGGACATAGAGTCAT 55
        |||||||
DB      1  GTCATCGATTGACCTAGTGCCTGGCCGGGCTTTGGCCGGCGGACATAGAGTCAT 55

```

RESULT 8
US-08-793-170-19/c
Sequence 19, Application US/08793170
Patent No. 5994128
GENERAL INFORMATION:
APPLICANT: FALLAUX et al.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
TITLE OF INVENTION: ADENOVIRUS TO BE USED IN GENE THERAPY
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESSES:
ADDRESSEE: RAE-VENTER LAW GROUP, P.C.
STREET: 260 SHERIDAN AVENUE, PO BOX 60039
CITY: PALO ALTO
STATE: CALIFORNIA
COUNTRY: USA
ZIP: 94306
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/793,170
FILING DATE: 25-MAR-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO 97/00326
FILING DATE: 14-JUN-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: EP 95201728.3
FILING DATE: 26-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: EP 95201611.1
FILING DATE: 15-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: RAE-VENTER, BARBARA
REGISTRATION NUMBER: 32,750
REFERENCE/DOCKET NUMBER: INGE.002.00US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650)328-4400
TELEFAX: (650)328-3377
TELEX: N/A
INFORMATION FOR SEQ ID NO: 19:
SEQUENCE CHARACTERISTICS:
LENGTH: 55 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
US-08-793-170-19

Query Match 92.7%; Score 51; DB 2; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.4e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 ATCGATTGACCTAGTGGCCGCGGCTTTGCCCGGCGGCGACATAGTCAAT 55
|||||
DB 55 ATCGATTGACCTAGTGGCCGCGGCTTTGCCCGGCGGCGACATAGTCAAT 5

RESULT 9
US-08-892-873-19/c
Sequence 19, Application US/08892873
Patent No. 6033908
GENERAL INFORMATION:
APPLICANT: FALLAUX et al.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
TITLE OF INVENTION: ADENOVIRUS TO BE USED IN GENE THERAPY
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESSES:
ADDRESSEE: RAE-VENTER LAW GROUP, P.C.
STREET: 260 SHERIDAN AVENUE, PO BOX 60039

CITY: PALO ALTO
STATE: CALIFORNIA
COUNTRY: USA
ZIP: 94306
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/892,873
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/793,170
FILING DATE: 25-MAR-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO 97/00326
FILING DATE: 14-JUN-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: EP 95201728.3
FILING DATE: 26-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: EP 95201611.1
FILING DATE: 15-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: RAE-VENTER, BARBARA
REGISTRATION NUMBER: 32,750
REFERENCE/DOCKET NUMBER: INGE.002.00US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650)328-4400
TELEFAX: (650)328-3377
TELEX: N/A
INFORMATION FOR SEQ ID NO: 19:
SEQUENCE CHARACTERISTICS:
LENGTH: 55 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
US-08-892-873-19

Query Match 92.7%; Score 51; DB 3; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.4e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 ATCGATTGACCTAGTGGCCGCGGCTTTGCCCGGCGGCGACATAGTCAAT 55
|||||
DB 55 ATCGATTGACCTAGTGGCCGCGGCTTTGCCCGGCGGCGACATAGTCAAT 5

RESULT 10
US-09-334-765A-19/c
Sequence 19, Application US/09334765A
Patent No. 6238893
GENERAL INFORMATION:
APPLICANT: Fallaux, Frits J.
APPLICANT: Hoebein, Robert C.
APPLICANT: Bouc, Abraham
APPLICANT: Valetto, Domenico
APPLICANT: Van der Eb, Alex J.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO BE USED
FILE REFERENCE: 3833.2US
CURRENT APPLICATION NUMBER: US/09/334,765A
CURRENT FILING DATE: 1999-06-16
PRIOR APPLICATION NUMBER: US 08/793,170
PRIOR FILING DATE: 1997-03-25
PRIOR APPLICATION NUMBER: PCT/NL96/00244
PRIOR FILING DATE: 1996-06-14
PRIOR APPLICATION NUMBER: EP 95201728.3
PRIOR FILING DATE: 1995-06-26
PRIOR APPLICATION NUMBER: EP 95201611.1

```
; PRIOR FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 19
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/clal
US-09-334-765A-19
```

```
Query Match          92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.4e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 5 ATCGATTGACCTAGTGGCCGGGCGGCTTTGGCCGGGGGCGGCACTAGGTCAAT 55
Db 55 ATCGATTGACCTAGTGGCCGGGCGGCTTTGGCCGGGGGCGGCACTAGGTCAAT 5
```

RESULT 11

```
; Sequence 19, Application US/09356575E
; Patent No. 6265212
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits
; APPLICANT: Hoebein, Robert
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: van der Eb, Alex
; APPLICANT: Schouten, Govert
; TITLE OF INVENTION: PACKAGING SYSTEMS
; FILE REFERENCE: 2578-393505
; CURRENT APPLICATION NUMBER: US/09/356,575E
; CURRENT FILING DATE: 1999-07-19
; PRIOR APPLICATION NUMBER: US 08/793,170
; PRIOR FILING DATE: 1997-03-25
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 19
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; OTHER INFORMATION: Derived from Adenovirus
US-09-356-575E-19
```

```
Query Match          92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.4e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 5 ATCGATTGACCTAGTGGCCGGGCGGCTTTGGCCGGGGGCGGCACTAGGTCAAT 55
Db 55 ATCGATTGACCTAGTGGCCGGGCGGCTTTGGCCGGGGGCGGCACTAGGTCAAT 5
```

RESULT 12

```
; Sequence 19, Application US/09333820A
; Patent No. 6306652
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoebein, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
```

```
; APPLICANT: Van der Eb, Alex J.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO BE USED
```

```
; TITLE OF INVENTION: GENE THERAPY
; FILE REFERENCE: 3833.1US
; CURRENT APPLICATION NUMBER: US/09/333,820A
; CURRENT FILING DATE: 1999-06-15
; EARLIER APPLICATION NUMBER: US 08/793,170
; EARLIER FILING DATE: 1997-03-25
; EARLIER APPLICATION NUMBER: PCT/NL96/00244
; EARLIER FILING DATE: 1996-06-14
; EARLIER APPLICATION NUMBER: EP 95201611.1
; EARLIER FILING DATE: 1995-06-26
; EARLIER APPLICATION NUMBER: EP 95201728.3
; EARLIER FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 19
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/clal
US-09-333-820-19
```

```
Query Match          92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.4e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 5 ATCGATTGACCTAGTGGCCGGGCGGCTTTGGCCGGGGGCGGCACTAGGTCAAT 55
Db 55 ATCGATTGACCTAGTGGCCGGGCGGCTTTGGCCGGGGGCGGCACTAGGTCAAT 5
```

RESULT 13

```
; Sequence 41, Application US/09358036
; Patent No. 6340595
; GENERAL INFORMATION:
; APPLICANT: Vogels, Ronald
; APPLICANT: Bout, Abraham
; APPLICANT: van Es, Helmut
; APPLICANT: Schouten, Govert
; TITLE OF INVENTION: High Throughput Screening of Gene Function Using
; TITLE OF INVENTION: Adenoviral Libraries for Functional Genomics
; FILE REFERENCE: 2184108
; CURRENT APPLICATION NUMBER: US/09/358,036
; CURRENT FILING DATE: 1999-07-21
; EARLIER APPLICATION NUMBER: US 09/097,239
; EARLIER FILING DATE: 1995-07-25
; NUMBER OF SEQ ID NOS: 69
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 41
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: oligonucleotide
US-09-358-036-41
```

```
Query Match          92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.4e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 5 ATCGATTGACCTAGTGGCCGGGCGGCTTTGGCCGGGGGCGGCACTAGGTCAAT 55
Db 55 ATCGATTGACCTAGTGGCCGGGCGGCTTTGGCCGGGGGCGGCACTAGGTCAAT 5
```

RESULT 14

```
; Sequence 19, Application US/09333820A
; Patent No. 6306652
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoebein, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
```

```
; Sequence 41, Application US/09097239
; Patent No. 641376
; GENERAL INFORMATION:
; APPLICANT: VOGELS, RONALD,
; APPLICANT: BOUT, ABRAHAM,
; APPLICANT: VAN ES, HELMUTH HG,
; APPLICANT: SCHOUTEN, GOVERT
; TITLE OF INVENTION: HIGH THROUGHPUT SCREENING OF GENE
; TITLE OF INVENTION: FUNCTION USING ADENOVIRAL LIBRARIES FOR FUNCTIONAL
; NUMBER OF SEQUENCES: 47
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: RAE-VENTER LAW GROUP
; STREET: PO BOX 60039
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94306
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/097,239
; FILING DATE: 12-JUN-1998
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: RAE-VENTER, BARBARA
; REGISTRATION NUMBER: 32,750
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650)328-4400
; TELEFAX: (650)328-4477
; INFORMATION FOR SEQ ID NO: 41:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 55 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; US-09-097-239-41

Query Match          92.7%; Score 51; DB 4; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.4e-09;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 ATGATGACCTAGTGGCGCGGCTTTCGCCGGCGGCACTAGTCAAT 55
Db 55 ATGATGACCTAGTGGCGCGGCTTTCGCCGGCGGCACTAGTCAAT 5

RESULT 15
US-08-793-170-17/c
; Sequence 17, Application US/08793170
; Patent No. 5994128
; GENERAL INFORMATION:
; APPLICANT: FALLAUX et al.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT
; TITLE OF INVENTION: ADENOVIRUS TO BE USED IN GENE THERAPY
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: RAE-VENTER LAW GROUP, P.C.
; STREET: 260 SHERIDAN AVENUE, PO BOX 60039
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94306
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
```

```
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/793,170
; FILING DATE: 25-MAR-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO 97/00326
; FILING DATE: 14-JUN-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: EP 95201728.3
; FILING DATE: 26-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: EP 95201611.1
; FILING DATE: 15-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: RAE-VENTER, BARBARA
; REGISTRATION NUMBER: 32,750
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650)328-4400
; TELEFAX: (650)328-3377
; TELEX: N/A
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; US-08-793-170-17

Query Match          78.9%; Score 43.4; DB 2; Length 50;
Best Local Similarity 97.8%; Pred. No. 6.6e-07;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 11 TGACCTAGTGGCGCGGCTTTCGCCGGCGGCACTAGTCAAT 55
Db 49 TGACCTAGTGGCGCGGCTTTCGCCGGCGGCACTAGTCAAT 5
```

Search completed: December 27, 2002, 06:17:00
Job time : 34 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd

OM nucleic - nucleic search, using sw model

Run on: December 27, 2002, 00:33:34 ; Search time 170.5 Seconds
(without alignments)
726.451 Million cell updates/sec

```

Title:      US-09-918-029-20
Perfect score: 55
Sequence:   1  gtacatgatgacctagt.....ccggcgycactagtccaat 55

```

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

```
Searched:      2185239 segs, 1125999159 residues
Total number of hits satisfying chosen parameters: 4370478
```

```
Minimum DB seq length: 0
Maximum DB seq length: 20000000000
```

```
Post-processing: Minimum Match 0%
                  Maximum Match 100%
                  Listing first 45 summaries
```

Database : N Geneseq 101002:*

1:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1980.DAT:*
2:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1981.DAT:*
3:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1982.DAT:*
4:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1983.DAT:*
5:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1984.DAT:*
6:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1985.DAT:*
7:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1986.DAT:*
8:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1987.DAT:*
9:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1988.DAT:*
10:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1989.DAT:*
11:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1990.DAT:*
12:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1991.DAT:*
13:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1992.DAT:*
14:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1993.DAT:*
15:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1994.DAT:*
16:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1995.DAT:*
17:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1996.DAT:*
18:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1997.DAT:*
19:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1998.DAT:*
20:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA1999.DAT:*
21:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA2000.DAT:*
22:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA2001A.DAT:*
23:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
24:	/SID52/gcgdata/geneseq/geneseqn-emb1/NA2002.DAT:*

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	55	100.0	55	18	AA748647	Synthetic haipin o
2	55	100.0	55	21	AA259117	Oligonucleotide HP
3	55	100.0	55	24	ABK47033	Adenovirus vecto
4	51	92.7	55	18	AA748646	Synthetic haipin o
5	51	92.7	55	21	AA259116	Oligonucleotide HP
6	51	92.7	55	21	AA237959	Adenoviral constru
7	51	92.7	55	22	AAF30332	Oligonucleotide fd
8	51	92.7	55	24	ABK47032	Adenovirus vecto
9	43.4	78.9	50	18	AA748644	Synthetic haipin o

10	43.4	78.9	50	18	AA748645
11	43.4	78.9	50	21	AA37958
12	43.4	78.9	50	22	AA30231
13	43.4	78.9	50	24	ABK47030
14	43.4	78.9	50	24	ABK47031
15	43	78.2	54	21	AA37960
16	42.2	76.7	55	18	AA748647
17	42.2	76.7	55	18	AA748646
18	42.2	76.7	55	21	AA559116
19	42.2	76.7	55	21	AA559117
20	42.2	76.7	55	21	AA37959
21	42.2	76.7	55	22	AA30232
22	42.2	76.7	55	24	ABK47032
23	42.2	76.7	55	24	ABK47033
24	39.4	71.6	45	21	AA559131
25	39.4	71.6	45	21	AA37961
26	39.4	71.6	45	22	AA30234
27	39.4	71.6	45	24	ABK47038
28	38.6	70.2	50	18	AA748644
29	38.6	70.2	50	18	AA748645
30	38.6	70.2	50	21	AA37958
31	38.6	70.2	50	22	AA30231
32	38.6	70.2	50	24	ABK47030
33	38.6	70.2	50	24	ABK47031
34	34.6	62.9	45	21	AA529131
35	34.6	62.9	45	21	AA527961
36	34.6	62.9	45	22	AA30234
37	34.6	62.9	45	24	ABK47038
38	31.4	57.1	49	21	AA37957
39	30.2	54.9	54	21	AA37960
40	26.6	48.4	49	21	AA37957
41	23.8	43.3	849	24	ABN69129
42	23.8	43.3	894	22	AA382907
43	23.6	42.9	63	24	AB187600
44	23.6	42.9	145	14	AAQ41448
45	23.6	42.9	145	16	AA703855

ALIGNMENTS

RESULT 1
AAT48647
ID AAT48647 standard; DNA; 55 BP.
XX AAT48647;
AC
XX
DT 21-MAY-1997 (first entry)
DE
XX Synthetic haipin oligonucleotide HP/cle2.
XX
KW Gene therapy; vaccine; vector; adenovirus; packaging system;
KW haipin; PICU; ss.
XX
OS Synthetic.
XX
PN WO9700326-A1.
XX
PD 03-JAN-1997.
XX
PF 14-JUN-1996; 96WO-NL00244.
XX
PR 26-JUN-1995; 95EP-0201728.
PR 15-JUN-1995; 95EP-0201611.
XX
PA (INTR-) INTROGENE BV.
XX (UYLE-) RIJKSUNIV LEIDEN.
XX
PI Bout A, Fallaux FJ, Hoeben FC, Valerio D, Van Der Ebbad;
XX WPI; 1997-077531/07.
XX
PT New packaging cells and nucleic acids for recombinant adenovirus -

PF have no overlapping sequences, prevents homologous recombination;
 PF for use in gene therapy and vaccination
 XX
 PS Disclosure; Page 55; 88pp; English.
 CC Synthetic oligonucleotides HP/clal (AAT48646) and HP/claz (AAT48647)
 CC were used to generate a synthetic hairpin. They contain a ClaI
 CC recognition site to be used for hairpin formation. The
 CC oligonucleotides were annealed and ligated into plasmid pCMV-TK,
 CC at the adenovirus inverted terminal repeat, generating
 CC pAd-CMV-hcTK. This plasmid was co-transfected with ClaI-digested
 CC wild-type adenovirus 4 into 911 cells. A recombinant adenovirus
 CC in which the CMV-hcTK expression cassette replaced the EI sequences
 CC was isolated.
 SO Sequence 55 BP; 9 A; 17 C; 17 G; 12 T; 0 other;
 Query Match 100.0%; Score 55; DB 18; Length 55;
 Best Local Similarity 100.0%; Pred. No. 1.2e-10;
 Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 GTACATCATTTGACCTAGTCCGCCGGCTTTCGCCGGGCGGACATAGGTCAAT 55
 DB 1 GTACATCATTTGACCTAGTCCGCCGGCTTTCGCCGGGCGGACATAGGTCAAT 55
 RESULT 2
 ID AAS59117 standard; DNA; 55 BP.
 AC AAS59117;
 XX 11-APR-2000 (first entry)
 DT
 XX
 DE Oligonucleotide HP/claz for generating hairpin structure.
 XX
 KW Expressible nucleic acid library; gene expression; gene function;
 KM capillary formation; cell proliferation; hairpin structure; ss.
 XX
 OS Synthetic.
 XX
 PN MO9964582-A2.
 PD 16-DEC-1999.
 XX
 PF 11-JUN-1999; 99WO-NL00367.
 XX
 PR 12-JUN-1998; 98US-0097239.
 XX
 PA (IMPR-) INTRIGENE BV.
 PI Schouten G, Vogels R, Bout A, Van Es H;
 DR WPI; 2000-097536/08.
 XX
 PT New library of expressible nucleic acids, useful for high-throughput
 PT screening of gene function, especially for identifying therapeutic
 PT molecules -
 XX
 PS Example 3; Page 156; 223pp; English.
 CC The invention relates to a library of expressible nucleic acids (NA)
 CC which contains many compartments, each comprising at least one vehicle
 CC comprising at least one NA, the vehicle being capable of efficiently
 CC introducing a NA into a cell for expression. The library is useful for
 CC determining the function of one or more nucleic acids within the
 CC library, or to screen for an expressible nucleic acid with a particular
 CC desired function. It is especially useful for high throughput screening
 CC of gene function for functional genomics applications and for screening
 CC for nucleic acids with potential therapeutic value. Cell types
 CC appropriate for selection of a particular phenotype may be useful for
 CC capillary formation and cell proliferation. Oligonucleotides
 CC AAS59116-259117 were used to generate a hairpin structure in plasmid

CC pAd-CMV-hcTK. The hairpin structure was used to determine if it could be
 CC used to prime reverse strand synthesis on the displaced strand after
 CC replication initiation in the adenoviral inverted terminal repeat (ITR).
 CC
 XX
 PS Sequence 55 BP; 9 A; 17 C; 17 G; 12 T; 0 other;
 Query Match 100.0%; Score 55; DB 21; Length 55;
 Best Local Similarity 100.0%; Pred. No. 1.2e-10;
 Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 GTACATCATTTGACCTAGTCCGCCGGCTTTCGCCGGGCGGACATAGGTCAAT 55
 DB 1 GTACATCATTTGACCTAGTCCGCCGGCTTTCGCCGGGCGGACATAGGTCAAT 55
 RESULT 3
 ID ABR47033 standard; DNA; 55 BP.
 AC ABR47033;
 XX 05-JUN-2002 (first entry)
 DT
 XX
 DE Adenovirus vector pICLhae/haw hairpin linker sequence Hp/clal#2.
 XX
 KW Adenovirus vector library; ss; linker; high throughput screening;
 KM RCA; replication competent adenovirus.
 XX
 OS Synthetic.
 XX
 PN US6340595-B1.
 PD 22-JAN-2002.
 XX
 PF 21-JUL-1999; 99US-0358036.
 XX
 PR 12-JUN-1998; 98US-0097239.
 XX
 PA (GALA-) GALAPAGOS GENOMICS NV.
 PI Vogels R, Bout A, Van Es H, Schouten G;
 DR WPI; 2002-224926/28.
 XX
 PT Library of expressible nucleic acids, useful for determining nucleic
 PT acid function, comprises one or more adenoviral vectors capable of
 PT transfecting a host cell with the nucleic acid -
 XX
 PS Example 3; Column 85; 11pp; English.
 CC The invention relates to a library (I) of a multitude of unique
 CC expressible nucleic acids (NA), comprising a number of compartments
 CC (II), each consisting of one or more adenoviral vectors (III)
 CC comprising at least one unique NA of (I) in an aqueous medium, where
 CC (III) is capable of introducing the NA into a host cell (IV), is
 CC capable of expressing the product of the NA in (IV), and is deleted in
 CC a portion of the adenoviral genome necessary for replication. Also
 CC included is a method for producing the library. The library is useful
 CC for determining the function of at least one nucleic acid that is present.
 CC The library uses high throughput generation of recombinant adenoviral
 CC vector libraries containing one or more sample nucleic acids, followed by
 CC high throughput screening of the adenoviral vector libraries in a host to
 CC alter the phenotype of the host as a means of assigning a function to
 CC expression product(s) of the sample nucleic acids. The entire process
 CC lends itself to automation especially when implemented in a 96-well or
 CC other multi-well format. The high throughput screening, using a number of
 CC different in vitro assays, provides a means of efficiently obtaining
 CC functional information in a relatively short period of time. The
 CC member(s) of the recombinant adenoviral libraries that exhibit or induce
 CC a desired phenotype in a host in vitro or in situ are identified to
 CC reduce the libraries to a manageable number of recombinant adenoviral
 CC vectors or clones which can be tested in vitro in an animal model.
 CC Furthermore, the methods produce RCA-free adenoviral libraries. RCA

CC (replication competent adenovirus) contamination throughout the libraries
 CC could become a major obstacle, especially if libraries are continuously
 CC amplified for use in multiple screening programs. Additionally, a further
 CC advantage is minimisation of the number of steps involved in the process.
 CC There is no requirement for cloning each individual adenovirus before use
 CC in a high throughput screening program. Other systems include one or more
 CC steps in E. coli to achieve homologous recombination for the various
 CC adenoviral plasmids necessary for vector generation. The present
 CC sequence is a linker sequence used in the construction of the adenoviral
 CC vector library of the invention.

XX
 SQ Sequence 55 BP; 9 A; 17 C; 17 G; 12 T; 0 other;

Query Match 100.0%; Score 55; DB 24; Length 55;
 Best Local Similarity 100.0%; Pred. No. 1.2e-10;

Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTCATGCGATTGACCTAGTCCGCGGCTTGGCCGGCGGACACTAGTCAAT 55
 ||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 1 GTCATGCGATTGACCTAGTCCGCGGCTTGGCCGGCGGACACTAGTCAAT 55

RESULT 4

AA748646/C
 ID AA748646 standard; DNA; 55 BP.

XX
 AC AA748646;

XX
 DT 21-MAY-1997 (first entry)

XX
 DE Synthetic hairpin oligonucleotide HP/c1a1.

XX
 KW Gene therapy; vaccine; vector; adenovirus; packaging system;
 hairpin; pICL; ss.

XX
 OS Synthetic.

XX
 PN WO9700326-A1.

XX
 PD 03-JAN-1997.

XX
 PF 14-JUN-1996; 96WO-NL00244.

XX
 PR 26-JUN-1995; 95EP-0201728.

XX
 PR 15-JUN-1995; 95EP-0201611.

XX
 PA (INTR-) INTROGENE BV.

XX
 PA (UYLE-) RICKSONIV LEIDEN.

XX
 PI Bout A, Fallaux FJ, Hoebe RC, Valerio D, Van Der EBBALJ;
 WPI; 1997-077531/07.

XX
 DR New packaging cells and nucleic acids for recombinant adenovirus -
 PT have no overlapping sequences, prevents homologous recombination;
 PT for use in gene therapy and vaccination

XX
 PS Disclosure; Page 55; 88pp; English.

XX
 SQ Synthetic oligonucleotides HP/c1a1 (AA748646) and HP/c1a2 (AA748647)

CC were used to generate a synthetic hairpin. They contain a c1a1
 CC recognition site to be used for hairpin formation. The

CC oligonucleotides were annealed and ligated into plasmid pCMV.TK,
 CC at the adenovirus inverted terminal repeat, generating

CC PAD-CMV-hcTK. This plasmid was co-transfected with c1a1-digested
 CC wild-type adenovirus 4 into 911 cells. A recombinant adenovirus

CC in which the CMV-hcTK expression cassette replaced the E1 sequences
 CC was isolated.

XX
 SQ Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;

Query Match 92.7%; Score 51; DB 18; Length 55;
 Best Local Similarity 100.0%; Pred. No. 3.4e-09;

Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 ATCGATTGACCTAGTCCGCGGCTTGGCCGGCGGACACTAGTCAAT 55
 ||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 55 ATCGATTGACCTAGTCCGCGGCTTGGCCGGCGGACACTAGTCAAT 5

RESULT 5

AA259116/C
 ID AA259116 standard; DNA; 55 BP.

XX
 AC AA259116;

XX
 DT 11-APR-2000 (first entry)

XX
 DE Oligonucleotide HP/c1a1 for generating hairpin structure.

XX
 KW Expressible nucleic acid library; gene expression; gene function;
 capillary formation; cell proliferation; hairpin structure; ss.

XX
 OS Synthetic.

XX
 PN WO964582-A2.

XX
 PD 16-DEC-1999.

XX
 PF 11-JUN-1999; 99WO-NL00367.

XX
 PR 12-JUN-1998; 98US-0097239.

XX
 PA (INTR-) INTROGENE BV.

XX
 PI Schouten G, Vogels R, Bout A, Van Es H;
 WPI; 2000-097536/08.

XX
 DR New library of expressible nucleic acids, useful for high-throughput
 PT screening of gene function, especially for identifying therapeutic
 PT molecules

XX
 PS Example 3; Page 156; 223pp; English.

XX
 The invention relates to a library of expressible nucleic acids (NA)
 CC which contains many compartments, each comprising at least one vehicle
 CC comprising at least one NA, the vehicle being capable of efficiently
 CC introducing a NA into a cell for expression. The library is useful for
 CC determining the function of one or more nucleic acids within the
 CC library, or to screen for an expressible nucleic acid with a particular
 CC desired function. It is especially useful for high throughput screening
 CC of gene function for functional genomics applications and for screening
 CC for nucleic acids with potential therapeutic value. Cell types
 CC appropriate for selection of a particular phenotype may be useful for
 CC capillary formation and cell proliferation. Oligonucleotides

XX
 AA259116-259117 were used to generate a hairpin structure in plasmid
 CC PAD-CMV-hcTK. The hairpin structure was used to determine if it could be
 CC used to prime reverse strand synthesis on the displaced strand after
 CC replication initiation in the adenoviral inverted terminal repeat (ITR).

XX
 SQ Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;

Query Match 92.7%; Score 51; DB 21; Length 55;
 Best Local Similarity 100.0%; Pred. No. 3.4e-09;

Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 ATCGATTGACCTAGTCCGCGGCTTGGCCGGCGGACACTAGTCAAT 55
 ||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 55 ATCGATTGACCTAGTCCGCGGCTTGGCCGGCGGACACTAGTCAAT 5

XX
 RESULT 6

AA237959/C
 ID AA237959 standard; DNA; 55 BP.

XX

AC AA237959;
 XX
 DT 07-FEB-2000 (first entry)
 XX
 DE Adenoviral construct generating primer Hp/c1a1.
 XX
 XX Adenoviral vector; replication-defective; adenovirus; ITR; hepatitis;
 KW inverted terminal repeat; encapsulation signal; gene therapy; tumor;
 KW inherited disease; cystic fibrosis; Duchenne muscular dystrophy;
 KW hypercholesterolemia; blood clotting disorder; hemophilia; restenosis;
 KW autoimmune disease; rheumatoid arthritis; PCR primer; ss.
 XX
 OS Synthetic.
 XX
 PN W09955132-A2.
 XX
 PD 04-NOV-1999.
 XX
 PF 23-APR-1999; 99WO-NI00235.
 XX
 PR 24-APR-1998; 98US-0065752.
 XX
 PA (INTR-) INTROGENE BV.
 XX
 PI Vogels R, Bout A;
 XX
 DR WPI: 2000-023229/02.
 XX
 PT New recombinant adenovirus vectors, used particularly for gene therapy
 PT for treating inherited or acquired diseases
 XX
 PS Disclosure; Page 118; 161pp; English.
 XX
 CC The invention provides methods of producing recombinant adenoviral
 CC vectors (Adv's) for generating replication-defective adenoviruses.
 CC Generating an Adv comprises fusing 2 partially overlapping sequences
 CC nucleic acid molecules that are capable of combining with each other to
 CC allow the generation of a physically linked nucleic acid comprising at
 CC least 2 functional adenoviral inverted terminal repeats (ITRs), a
 CC functional encapsulation signal and a nucleic acid of interest. The
 CC products can be used for gene therapy for treating inherited diseases
 CC e.g. cystic fibrosis, Duchenne muscular dystrophy,
 CC hypercholesterolemia, blood clotting disorders (hemophilia) or acquired
 CC diseases such as tumors, hepatitis, (auto)immune diseases, restenosis, or
 CC rheumatoid arthritis. Sequences AA237954-960 represent primers used for
 CC PCR amplification of DNA fragments used for generation of adenoviral
 CC constructs of the invention.
 CC
 SO Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;
 Query Match 92.7%; Score 51; DB 21; Length 55;
 Best Local Similarity 100.0%; Pred. No. 3.4e-09;
 Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 5 ATCGATTGACCTAGTGGCGGCGGCTTTGCGCGGCGGACACTAGTCAAT 55
 DB 55 ATCGATTGACCTAGTGGCGGCGGCTTTGCGCGGCGGACACTAGTCAAT 5
 RESULT 7
 AAF30232/c
 ID AAF30232 standard; DNA; 55 BP.
 XX
 AC AAF30232;
 XX
 DT 30-APR-2001 (first entry)
 XX
 DE Oligonucleotide forming hairpin structure.
 XX
 KW Adenovirus; vector; gene therapy; packaging cell; hairpin; ds.
 XX
 OS Synthetic.

PH Key Location/Qualifiers
 FT misc-feature 1..4
 FT /*tag- a
 FT /note- "single-stranded 5' overhang"
 FT 55
 FT misc-feature
 FT /*tag- b
 FT /note- "single-stranded overhang on complementary
 strand of sequence 5'-GTAC-3'
 XX
 PN W0200105945-A2.
 XX
 PD 25-JAN-2001.
 XX
 PF 19-JUL-2000; 2000WO-EP07074.
 XX
 PR 19-JUL-1999; 99US-0356575.
 XX
 PA (INTR-) INTROGENE BV.
 XX
 PI Hoeben RC, Bout A, Valerio D, Van Der Eb AJ, Schouten G;
 PI Fallaux FJ;
 XX
 DR WPI: 2001-147334/15.
 XX
 PT Producing recombinant adenovirus for use in gene therapy, comprises
 PT culturing cells containing adenoviral nucleic acid having an
 PT encapsidating signal and inverted terminal repeat, and lacking
 PT overlapping sequences
 XX
 PS Example; Page 39; 97pp; English.
 XX
 CC The present sequence is that of an oligonucleotide formed from 2
 CC partially complementary oligonucleotides creating a hairpin
 CC structure. The oligonucleotide forms an ClaI recognition site
 CC when inserted into the ClaI site of plasmid pICL (see AAF30233).
 CC This was performed as part of an experiment to determine whether
 CC the hairpin could be used as a primer for reverse strand synthesis
 CC on the displaced strand after replication had started from the
 CC inverted terminal repeat (ITR) of the vector. In adenovirus
 CC infected cells, linear DNA fragments have on one terminus an
 CC adenovirus-derived ITR and at the other terminus a sequence that
 CC can anneal to the same strand, when present in single-stranded
 CC form, thereby generating a hairpin structure, and will be
 CC converted to structures with ITRs at both ends. The resulting DNA
 CC molecules will replicate by the same mechanism as the wild-type
 CC adenovirus genomes. The invention provides adenovirus vectors and
 CC packaging cell lines useful in the safe generation of EI-deleted
 CC recombinant adenovirus vectors for gene therapy applications.
 CC Packaging cells contain adenovirus nucleic acids having an
 CC encapsidating signal and ITR, but lack sequences that overlap with
 CC the vector, thereby preventing homologous recombination.
 CC
 SO Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;
 Query Match 92.7%; Score 51; DB 22; Length 55;
 Best Local Similarity 100.0%; Pred. No. 3.4e-09;
 Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 5 ATCGATTGACCTAGTGGCGGCGGCTTTGCGCGGCGGACACTAGTCAAT 55
 DB 55 ATCGATTGACCTAGTGGCGGCGGCTTTGCGCGGCGGACACTAGTCAAT 5
 RESULT 8
 ABK47032/c
 ID ABK47032 standard; DNA; 55 BP.
 XX
 AC ABK47032;
 XX
 DT 05-JUN-2002 (first entry)
 XX
 DE Adenovirus vectro pICuhac/haw hairpin linker sequence Hp/c1a1#1.

KM Adenovirus vector library; ss; linker; high throughput screening;
 KM RCA; replication competent adenovirus.
 XX
 OS Synthetic.
 XX
 PN US6340595-B1.
 XX
 PD 22-JAN-2002.
 XX
 PF 21-JUL-1999; 99US-0358036.
 XX
 PR 12-JUN-1998; 98US-0097239.
 XX
 PA (GALA-) GALAPAGOS GENOMICS NV.
 XX
 PI Vogels R, Bout A, Van Es H, Schouten G;
 DR WPI; 2002-224926/28.
 XX
 PT Library of expressible nucleic acids, useful for determining nucleic
 PT acid function, comprises one or more adenoviral vectors capable of
 PT transfecting a host cell with the nucleic acid -
 XX
 PS Example 3; Column 85; 11pp; English.
 XX
 CC The invention relates to a library (I) of a multitude of unique
 CC expressible nucleic acids (NA), comprises a number of compartments
 CC (II), each consisting of one or more adenoviral vectors (III)
 CC comprising at least one unique NA of (I) in an aqueous medium, where
 CC (III) is capable of introducing the NA into a host cell (IV), is
 CC capable of expressing the product of the NA in (IV), and is deleted in
 CC a portion of the adenoviral genome necessary for replication. Also
 CC included is a method for producing the library. The library is useful for
 CC determining the function of at least one nucleic acid that is present.
 CC The library uses high throughput generation of recombinant adenoviral
 CC vector libraries containing one or more sample nucleic acids, followed by
 CC high throughput screening of the adenoviral vector libraries in a host to
 CC alter the phenotype of the host as a means of assigning a function to
 CC expression product(s) of the sample nucleic acids. The entire process
 CC lends itself to automation especially when implemented in a 96-well or
 CC other multi-well format. The high throughput screening, using a number of
 CC different in vitro assays, provides a means of efficiently obtaining
 CC functional information in a relatively short period of time. The
 CC member(s) of the recombinant adenoviral libraries that exhibit or induce
 CC a desired phenotype in a host in vitro or in situ are identified to
 CC reduce the libraries to a manageable number of recombinant adenoviral
 CC vectors or clones which can be tested in vitro in an animal model.
 CC Furthermore, the methods produce RCA-free adenoviral libraries. RCA
 CC (replication competent adenovirus) contamination throughout the libraries
 CC could become a major obstacle, especially if libraries are continuously
 CC amplified for use in multiple screening programs. Additionally, a further
 CC advantage is minimisation of the number of steps involved in the process.
 CC There is no requirement for cloning each individual adenovirus before use
 CC in a high throughput screening program. Other systems include one or more
 CC steps in E. coli to achieve homologous recombination for the various
 CC adenoviral plasmids necessary for vector generation. The present
 CC sequence is a linker sequence used in the construction of the adenoviral
 CC vector library of the invention.
 XX
 SQ Sequence 55 BP; 12 A; 17 C; 17 G; 9 T; 0 other;
 XX
 Query Match 92.7%; Score 51; DB 24; Length 55;
 Best Local Similarity 100.0%; Pred. No. 3.4e-09;
 Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 QY 5 ATCGATTGACCTAGTGGCGCCGGGCTTTGGCCGGGCGGCACTAGTCAAT 55
 ||||||||||||||||||||||||||||||||||||||||||||||||
 DB 55 ATCGATTGACCTAGTGGCGCCGGGCTTTGGCCGGGCGGCACTAGTCAAT 5

XX
 AC AAT48644;
 XX
 DT 21-MAY-1997 (first entry)
 XX
 DE Synthetic hairpin oligonucleotide HP/asp1.
 XX
 KM Gene therapy; vaccine; vector; adenovirus; packaging system;
 KM hairpin; PICL; ss.
 XX
 OS Synthetic.
 XX
 PN WO9700326-A1.
 XX
 PD 03-JAN-1997.
 XX
 PF 14-JUN-1996; 96WO-NL00244.
 XX
 PR 26-JUN-1995; 95EP-0201728.
 PR 15-JUN-1995; 95EP-0201611.
 XX
 PA (INTR-) INTRIGENE BV.
 PA (UYLE-) RIJSDUWIV LEIDEN.
 XX
 PI Bout A, Fallaux FJ, Hoeven RC, Valerio D, Van Der EBBALJ;
 DR WPI; 1997-077531/07.
 XX
 PT New packaging cells and nucleic acids for recombinant adenovirus -
 PT have no overlapping sequences, prevents homologous recombination;
 PT for use in gene therapy and vaccination
 XX
 PS Disclosure; Page 55; 88pp; English.
 XX
 CC Synthetic oligonucleotides HP/asp1 (AAT48644) and HP/asp2 (AAT48645)
 CC were used to generate a synthetic hairpin, recreating an Asp718
 CC site at one of the terminal if inserted in the Asp718 site of
 CC adenovirus minimal vector PICL (see also AAT48630). Insertion of
 CC the oligonucleotides into PICL generated clone PICLinc (correct
 CC orientation) and PICLinc (reverse, non-functional orientation).
 CC The constructs were used to demonstrate the competence of a
 CC synthetic DNA sequence, that is capable of forming a hairpin
 CC structure, to serve as a primer for reverse strand synthesis in
 CC the generation of double-stranded DNA molecules in cells that
 CC contain and express adenovirus genes.
 CC
 SQ Sequence 50 BP; 10 A; 17 C; 17 G; 6 T; 0 other;
 XX
 Query Match 78.9%; Score 43.4; DB 18; Length 50;
 Best Local Similarity 97.8%; Pred. No. 1.8e-06;
 Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 XX
 QY 11 TGACCTAGTGGCGCCGGGCTTTGCCGGGCGGCACTAGTCAAT 55
 ||||||||||||||||||||||||||||||||||||||||||||||||
 DB 49 TGACCTAGTGGCGCCGGGCTTTGCCGGGCGGCACTAGTCAAT 5

RESULT 9
 AAT48644/c
 ID AAT48644 standard; DNA: 50 BP.

RESULT 10
 AAT48645
 ID AAT48645 standard; DNA: 50 BP.

```

XX 03-JAN-1997.
PD
DR
XX 14-JUN-1996; 96WO-NL00244.
PF
XX 26-JUN-1995; 95EP-0201728.
PR 15-JUN-1995; 95EP-0201611.
XX
PA (INTR-) INTROGENE BV.
PA (UYLE-) RIJCKSUNIV LEIDEN.
XX
PI Bout A, Fallaux FJ, Hoeben RC, Valerio D, Van Der EBBALJ;
DR WPI; 1997-077531/07.
XX
XX New packaging cells and nucleic acids for recombinant adenovirus -
PT have no overlapping sequences, prevents homologous recombination;
PT for use in gene therapy and vaccination
XX
PS Disclosure; Page 55; 88pp; English.
XX
XX Synthetic oligonucleotides Hp/asp1 (AAAT48644) and Hp/asp2 (AAAT48645)
CC were used to generate a synthetic hairpin, recreating an Asp718
CC site at one of the termini. If inserted in the Asp718 site of
CC adenovirus minimal vector pICL (see also AAT48630), insertion of
CC the oligonucleotides into pICL generated clone pICLhac (correct
CC orientation) and pICLhac (reverse, non-functional orientation).
CC The constructs were used to demonstrate the competence of a
CC synthetic DNA sequence, that is capable of forming a hairpin
CC structure, to serve as a primer for reverse strand synthesis in
CC the generation of double-stranded DNA molecules in cells that
CC contain and express adenovirus genes.
XX
SQ Sequence 50 BP; 6 A; 17 C; 17 G; 10 T; 0 other;
XX
Query Match 78.9%; Score 43.4; DB 18; Length 50;
Best Local Similarity 97.8%; Pred. No. 1.8e-06;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY 11 TGACCTAGTGC CGCCGCGGCTTTGCCGCGGCGCACTAGTCAAT 55
DB 6 TGACCTAGTGC CGCCGCGGCTTTGCCGCGGCGCACTAGTCAAGT 50
RESULT 11
AAZ37958
ID AAZ37958 standard; DNA; 50 BP.
XX
AC AAZ37958;
XX
DT 07-FEB-2000 (first entry)
XX
DE Adenoviral construct generating primer Hp/asp2.
XX
KW Adenoviral vector; replication-defective; adenovirus; ITR; hepatitis;
KW inverted terminal repeat; encapsulation signal; gene therapy; tumor;
KW inherited disease; cystic fibrosis; Duchenne molecular dystrophy;
KW hypercholesterolemia; blood clotting disorder; hemophilia; restenosis;
KW autoimmune disease; rheumatoid arthritis; PCR primer; ss.
XX
OS Synthetic.
XX
PN W09955132-A2.
XX
PD 04-NOV-1999.
XX
PF 23-APR-1999; 99WO-NL00235.
XX
PR 24-APR-1998; 98US-0065752.
XX
PA (INTR-) INTROGENE BV.
XX
PI Vogels R, Bout A;

```

```

XX WPI; 2000-023229/02.
DR
XX
XX New recombinant adenovirus vectors, used particularly for gene therapy
PT for treating inherited or acquired diseases -
XX
PS Disclosure; Page 118; 161pp; English.
XX
XX The invention provides methods of producing recombinant adenoviral
CC vectors (Adv's) for generating replication-defective adenoviruses.
CC Generating an Adv comprises fusing 2 partially overlapping sequences
CC nucleic acid molecules that are capable of combining with each other to
CC allow the generation of a physically linked nucleic acid comprising at
CC least 2 functional adenoviral inverted terminal repeats (ITRs), a
CC functional encapsulation signal and a nucleic acid of interest. The
CC products can be used for gene therapy for treating inherited diseases
CC e.g. cystic fibrosis, Duchenne molecular dystrophy,
CC hypercholesterolemia, blood clotting disorders (hemophilia) or acquired
CC diseases such as tumors, hepatitis, (auto)immune diseases, restenosis, or
CC rheumatoid arthritis. Sequences AAZ37954-960 represent primers used for
CC PCR amplification of DNA fragments used for generation of adenoviral
CC constructs of the invention.
XX
SQ Sequence 50 BP; 6 A; 17 C; 17 G; 10 T; 0 other;
XX
Query Match 78.9%; Score 43.4; DB 21; Length 50;
Best Local Similarity 97.8%; Pred. No. 1.8e-06;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY 11 TGACCTAGTGC CGCCGCGGCTTTGCCGCGGCGCACTAGTCAAT 55
DB 6 TGACCTAGTGC CGCCGCGGCTTTGCCGCGGCGCACTAGTCAAGT 50
RESULT 12
AAF30231/c
ID AAF30231 standard; DNA; 50 BP.
XX
AC AAF30231;
XX
DT 30-APR-2001 (first entry)
XX
DE Oligonucleotide forming hairpin structure.
XX
KW Adenovirus; vector; gene therapy; packaging cell; hairpin; ds.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT /*tag- a 1..4
FT misc-feature /note= "single-stranded 5' overhang"
FT misc-feature 50
FT /*tag- b
FT /*tag- b "single-stranded overhang on complementary
FT strand of sequence 5'-GTAC-3'"
XX
PN W0200105945-A2.
XX
PD 25-JAN-2001.
XX
XX 19-JUL-2000; 2000WO-EP07074.
XX
PF 19-JUL-1999; 99US-0356575.
XX
PR
XX
PA (INTR-) INTROGENE BV.
XX
PI Hoeben RC, Bout A, Valerio D, Van Der Eb AJ, Schouten G;
XX
PI Fallaux FJ;
XX
DR WPI; 2001-147334/15.
XX
PT Producing recombinant adenovirus for use in gene therapy, comprises

```

PT culturing cells containing adenoviral nucleic acid having an
PT encapsulating signal and inverted terminal repeat, and lacking
PT overlapping sequences -

XX Example: Page 39; 97pp; English.

CC The present sequence is that of an oligonucleotide formed from 2
CC partially complementary oligonucleotides creating a hairpin
CC structure. The oligonucleotide forms an Asp718 recognition site
CC when inserted into the Asp718 site of plasmid pICL (see AAF0233).
CC This was performed as part of an experiment to determine whether
CC the hairpin could be used as a primer for reverse strand synthesis
CC on the displaced strand after replication had started from the
CC inverted terminal repeat (ITR) of the vector. In adenovirus
CC infected cells, linear DNA fragments have on one terminus an
CC adenovirus-derived ITR and at the other terminus a sequence that
CC can anneal to the same strand, when present in single-stranded
CC form, thereby generating a hairpin structure, and will be
CC converted to structures with ITRs at both ends. The resulting DNA
CC molecules will replicate by the same mechanism as the wild-type
CC adenovirus genomes. The invention provides adenovirus vectors and
CC packaging cell lines useful in the safe generation of EI-deleted
CC recombinant adenovirus vectors for gene therapy applications.
CC Packaging cells contain adenovirus nucleic acids having an
CC encapsulating signal and ITR, but lack sequences that overlap with
CC the vector, thereby preventing homologous recombination.

XX Sequence 50 BP; 10 A; 17 C; 17 G; 6 T; 0 other;

Query Match 78.9%; Score 43.4; DB 22; Length 50;
Best Local Similarity 97.8%; Pred. No. 1.8e-06;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 11 TGACCTAGTGGCCGCCGGCTTGGCCGGCGGCACTAGTCAAT 55
DB 49 TGACCTAGTGGCCGCCGGCTTGGCCGGCGGCACTAGTCACT 5

RESULT 13

ID ABR47030 standard; DNA: 50 BP.

XX ABR47030;

DT 05-JUN-2002 (first entry)

DE Adenovirus vectro pICLhac/haw linker sequence Hp/asp1#1.

XX Adenovirus vector library; ss; linker; high throughput screening;

KW RCA; replication competent adenovirus.

XX Synthetic.

PN US6340595-B1.

PD 22-JAN-2002.

PF 21-JUL-1999; 99US-0358036.

PR 12-JUN-1998; 98US-0097239.

XX (GALA-) GALAPAGOS GENOMICS NV.

PI Vogels R, Bout A, Van Es H, Schouten G;

DR WPI; 2002-224926/28.

XX Library of expressible nucleic acids, useful for determining nucleic
PT acid function, comprises one or more adenoviral vectors capable of
PT transfecting a host cell with the nucleic acid -
XX Example 3; Column 84; 11pp; English.

CC The invention relates to a library (I) of a multitude of unique
CC expressible nucleic acids (NA), comprises a number of compartments
CC (II), each consisting of one or more adenoviral vectors (III)
CC comprising at least one unique NA of (I) in an aqueous medium, where
CC (III) is capable of introducing the NA into a host cell (IV), is
CC capable of expressing the product of the NA in (IV), and is deleted in
CC a portion of the adenoviral genome necessary for replication. Also
CC included is a method for producing the library. The library is useful for
CC determining the function of at least one nucleic acid that is present.
CC The library uses high throughput generation of recombinant adenoviral
CC vector libraries containing one or more sample nucleic acids, followed by
CC high throughput screening of the adenoviral vector libraries in a host to
CC alter the phenotype of the host as a means of assigning a function to
CC expression product(s) of the sample nucleic acids. The entire process
CC lends itself to automation especially when implemented in a 96-well or
CC other multi-well format. The high throughput screening, using a number of
CC different in vitro assays, provides a means of efficiently obtaining
CC functional information in a relatively short period of time. The
CC member(s) of the recombinant adenoviral libraries that exhibit or induce
CC a desired phenotype in a host in vitro or in situ are identified to
CC reduce the libraries to a manageable number of recombinant adenoviral
CC vectors or clones which can be tested in vitro in an animal model.
CC Furthermore, the methods produce RCA-free adenoviral libraries. RCA
CC (replication competent adenovirus) contamination throughout the libraries
CC could become a major obstacle, especially if libraries are continuously
CC amplified for use in multiple screening programs. Additionally, a further
CC advantage is minimisation of the number of steps involved in the process.
CC There is no requirement for cloning each individual adenovirus before use
CC in a high throughput screening program. Other systems include one or more
CC steps in E. coli to achieve homologous recombination for the various
CC adenoviral plasmids necessary for vector generation. The present
CC sequence is a linker sequence used in the construction of the adenoviral
CC vector library of the invention.

XX Sequence 50 BP; 10 A; 17 C; 17 G; 6 T; 0 other;

Query Match 78.9%; Score 43.4; DB 24; Length 50;
Best Local Similarity 97.8%; Pred. No. 1.8e-06;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 11 TGACCTAGTGGCCGCCGGCTTGGCCGGCGGCACTAGTCAAT 55
DB 49 TGACCTAGTGGCCGCCGGCTTGGCCGGCGGCACTAGTCACT 5

RESULT 14

ID ABR47031 standard; DNA: 50 BP.

XX ABR47031;

DT 05-JUN-2002 (first entry)

DE Adenovirus vectro pICLhac/haw linker sequence Hp/asp1#2.

XX Adenovirus vector library; ss; linker; high throughput screening;

KW RCA; replication competent adenovirus.

XX Synthetic.

PN US6340595-B1.

PD 22-JAN-2002.

PF 21-JUL-1999; 99US-0358036.

PR 12-JUN-1998; 98US-0097239.

XX (GALA-) GALAPAGOS GENOMICS NV.

PI Vogels R, Bout A, Van Es H, Schouten G;

DR WPI; 2002-224926/28.

```

XX Library of expressible nucleic acids, useful for determining nucleic
PT acid function, comprises one or more adenoviral vectors capable of
PT transfecting a host cell with the nucleic acid
XX
PS Example 3; Column 84; 11pp; English.
XX
CC The invention relates to a library (I) of a multitude of unique
CC expressible nucleic acids (NA), comprises a number of compartments
CC (II), each consisting of one or more adenoviral vectors (III)
CC comprising at least one unique NA of (I) in an aqueous medium, where
CC (III) is capable of introducing the NA into a host cell (IV), is
CC capable of expressing the product of the NA in (IV), and is deleted in
CC a portion of the adenoviral genome necessary for replication. Also
CC included is a method for producing the library. The library is useful for
CC determining the function of at least one nucleic acid that is present.
CC The library uses high throughput generation of recombinant adenoviral
CC vector libraries containing one or more sample nucleic acids, followed by
CC high throughput screening of the adenoviral vector libraries in a host to
CC alter the phenotype of the host as a means of assigning a function to
CC expression product(s) of the sample nucleic acids. The entire process
CC lends itself to automation especially when implemented in a 96-well or
CC other multi-well format. The high throughput screening, using a number of
CC different in vitro assays, provides a means of efficiently obtaining
CC functional information in a relatively short period of time. The
CC member(s) of the recombinant adenoviral libraries that exhibit or induce
CC a desired phenotype in a host in vitro or in situ are identified to
CC reduce the libraries to a manageable number of recombinant adenoviral
CC vectors or clones which can be tested in vitro in an animal model.
CC Furthermore, the methods produce RCA-free adenoviral libraries. RCA
CC (replication competent adenovirus) contamination throughout the libraries
CC could become a major obstacle, especially if libraries are continuously
CC amplified for use in multiple screening programs. Additionally, a further
CC advantage is minimisation of the number of steps involved in the process.
CC There is no requirement for cloning each individual adenovirus before use
CC in a high throughput screening program. Other systems include one or more
CC steps in E. coli to achieve homologous recombination for the various
CC adenoviral plasmids necessary for vector generation. The present
CC sequence is a linker sequence used in the construction of the adenoviral
CC vector library of the invention.
XX
SQ Sequence 50 BP; 6 A; 17 C; 17 G; 10 T; 0 other;
Query Match 78.9%; Score 43.4; DB 24; Length 50;
Best Local Similarity 97.8%; Pred. No. 1.8e-06;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 11 TGACCTAGTGGCGCGCTTTCGCCGCGGCACTAGTCAAT 55
DB 6 TGACCTAGTGGCGCGCTTTCGCCGCGGCACTAGTCAAT 50
RESUME 15
AA237960
ID AA237960 standard; DNA; 54 BP.
AC AA237960;
XX
XX 07-FEB-2000 (first entry)
XX
XX Adenoviral construct generating primer Hp/cia2.
XX
XX Adenoviral vector; replication-defective; adenovirus; ITR; hepatitis;
XX inverted terminal repeat; encapsulation signal; gene therapy; tumor;
XX inherited disease; cystic fibrosis; Duchenne molecular dystrophy;
XX hypercholesterolemia; blood clotting disorder; hemophilia; restenosis;
XX autoimmune disease; rheumatoid arthritis; PCR primer; ss.
XX
XX Synthetic.
XX
XX WO9955132-A2.
XX
XX 04-NOV-1999.

```

```

XX
XX 23-APR-1999; 99WO-NL00235.
XX
XX 24-APR-1998; 98US-0065752.
XX
XX (INTR-) INTROGENE BV.
XX
XX
XX Vogels R, Bout A;
XX
XX WPI; 2000-023229/02.
XX
XX
XX New recombinant adenovirus vectors, used particularly for gene therapy
XX for treating inherited or acquired diseases
XX
XX Disclosure; Page 118; 161pp; English.
XX
XX The invention provides methods of producing recombinant adenoviral
XX vectors (Adv's) for generating replication-defective adenoviruses.
XX generating an Adv comprises fusing 2 partially overlapping sequences
XX nucleic acid molecules that are capable of combining with each other to
XX allow the generation of a physically linked nucleic acid comprising at
XX least 2 functional adenoviral inverted terminal repeats (ITRs), a
XX functional encapsulation signal and a nucleic acid of interest. The
XX products can be used for gene therapy for treating inherited diseases
XX e.g. cystic fibrosis, Duchenne molecular dystrophy,
XX hypercholesterolemia, blood clotting disorders (hemophilia) or acquired
XX diseases such as tumors, hepatitis, (auto)immune diseases, restenosis, or
XX rheumatoid arthritis. Sequences AA237954-960 represent primers used for
XX PCR amplification of DNA fragments used for generation of adenoviral
XX constructs of the invention.
XX
SQ Sequence 54 BP; 9 A; 16 C; 17 G; 12 T; 0 other;
Query Match 78.2%; Score 43; DB 21; Length 54;
Best Local Similarity 98.2%; Pred. No. 2.6e-06;
Matches 54; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
QY 1 GTACATGATGTGACCTAGTGGCGCGGCTTTCGCCGCGGCACTAGTCAAT 55
DB 1 GTACATGATGTGACCTAGTGGCGCGGCTTTCGCCGCGGCACTAGTCAAT 54
Search completed: December 27, 2002, 04:46:16
Job time : 171.5 secs

```


GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 27, 2002, 04:41:15 ; Search time 36 Seconds
(without alignments)
620.636 Million cell updates/sec

Title: US-09-918-029-20

Perfect score: 1 gtacatcgattgactagtagtgcacgagcgcagctagctaat 55

Sequence: IDENTITY_NUC

Scoring table: Gapop 10.0 , Gapext 1.0

Searched: 363474 seqs, 203117208 residues

Total number of hits satisfying chosen parameters: 726948

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications_NA:*

1: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq:*
2: /cgn2_6/ptodata/2/pubpna/PCT_NEW_PUB.seq:*
3: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq:*
4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq:*
5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq:*
6: /cgn2_6/ptodata/2/pubpna/PCTUS_PUBCOMB.seq:*
7: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq:*
8: /cgn2_6/ptodata/2/pubpna/US08_PUBCOMB.seq:*
9: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq:*
10: /cgn2_6/ptodata/2/pubpna/US09_PUBCOMB.seq:*
11: /cgn2_6/ptodata/2/pubpna/US10_NEW_PUB.seq:*
12: /cgn2_6/ptodata/2/pubpna/US10_PUBCOMB.seq:*
13: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq:*
14: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	55	100.0	55	9	US-10-125-751-20
2	55	100.0	55	9	US-09-912-552-20
3	55	100.0	55	10	US-09-918-029-20
4	55	100.0	55	12	US-10-038-271-20
5	51	92.7	55	9	US-10-125-751-19
6	51	92.7	55	9	US-09-912-552-19
7	51	92.7	55	10	US-09-918-029-19
8	51	92.7	55	10	US-09-900-062-45
9	51	92.7	55	12	US-10-038-271-19
10	43.4	78.9	50	9	US-10-125-751-18
11	43.4	78.9	50	9	US-10-125-751-17
12	43.4	78.9	50	9	US-09-912-552-17
13	43.4	78.9	50	9	US-09-918-029-17
14	43.4	78.9	50	10	US-09-918-029-18
15	43.4	78.9	50	10	US-09-900-062-44
16	43.4	78.9	50	12	US-10-038-271-17
17	43.4	78.9	50	12	US-10-038-271-18
18	43.4	78.9	50	12	US-10-038-271-17
19	43	78.2	54	10	US-09-900-062-46

20	42.2	76.7	55	9	US-10-125-751-19	Sequence 19, Appl
21	42.2	76.7	55	9	US-10-125-751-20	Sequence 20, Appl
22	42.2	76.7	55	9	US-09-912-552-19	Sequence 19, Appl
23	42.2	76.7	55	9	US-09-912-552-20	Sequence 20, Appl
24	42.2	76.7	55	10	US-09-918-029-19	Sequence 19, Appl
25	42.2	76.7	55	10	US-09-918-029-20	Sequence 20, Appl
26	42.2	76.7	55	12	US-09-900-062-45	Sequence 45, Appl
27	42.2	76.7	55	12	US-10-038-271-19	Sequence 19, Appl
28	42.2	76.7	55	12	US-10-038-271-20	Sequence 20, Appl
29	39.4	71.6	45	9	US-10-125-751-22	Sequence 22, Appl
30	39.4	71.6	45	10	US-09-918-029-22	Sequence 22, Appl
31	39.4	71.6	45	10	US-09-900-062-47	Sequence 47, Appl
32	39.4	71.6	45	12	US-10-038-271-22	Sequence 22, Appl
33	38.6	70.2	50	9	US-10-125-751-17	Sequence 17, Appl
34	38.6	70.2	50	9	US-10-125-751-18	Sequence 18, Appl
35	38.6	70.2	50	9	US-09-912-552-17	Sequence 17, Appl
36	38.6	70.2	50	9	US-09-912-552-18	Sequence 18, Appl
37	38.6	70.2	50	10	US-09-918-029-17	Sequence 17, Appl
38	38.6	70.2	50	10	US-09-918-029-18	Sequence 18, Appl
39	38.6	70.2	50	10	US-09-900-062-44	Sequence 44, Appl
40	38.6	70.2	50	12	US-10-038-271-17	Sequence 17, Appl
41	38.6	70.2	45	9	US-10-038-271-18	Sequence 18, Appl
42	36.2	65.8	45	9	US-09-912-552-22	Sequence 22, Appl
43	34.6	62.9	45	9	US-10-125-751-22	Sequence 22, Appl
44	34.6	62.9	45	10	US-09-918-029-22	Sequence 22, Appl
45	34.6	62.9	45	10	US-09-900-062-47	Sequence 47, Appl

ALIGNMENTS

RESULT 1
US-10-125-751-20
Sequence 20, Application US/10125751
Patent No. US20020173039A1
GENERAL INFORMATION:
APPLICANT: Fallaux, Fris J.
APPLICANT: Hoeber, Robert C.
APPLICANT: Bout, Abraham
APPLICANT: Valerio, Domenico
APPLICANT: Van der Eb, Alex J.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO
TITLE OF INVENTION: BE USED
FILE REFERENCE: 3833.2US
CURRENT APPLICATION NUMBER: US/10/125, 751
CURRENT FILING DATE: 2002-04-18
PRIOR APPLICATION NUMBER: 09/506, 548
PRIOR FILING DATE: 2000-02-16
PRIOR APPLICATION NUMBER: PCT/NL96/00244
PRIOR FILING DATE: 1996-06-14
PRIOR APPLICATION NUMBER: EP 95201728.3
PRIOR FILING DATE: 1995-06-26
PRIOR APPLICATION NUMBER: EP 95201611.1
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Corel Wordperfect 8.0
SEQ ID NO 20
LENGTH: 55
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
NAME/KEY:
LOCATION:
OTHER INFORMATION: Description of Artificial Sequence: primer HP/claz
US-10-125-751-20

Query Match 100.0%; Score 55; DB 9; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.5e-12;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
07 1 GTACATGATTGACTATGTCGCCGCCGCGCTTGCCTCCGCGGACACAGATGAT 55
|||||

```
Db      1  GTACATGATGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACACTAGTCAAT 55
RESULT 2
US-09-912-552-20
; Sequence 20, Application US/09912552
; Publication No. US20020187553A1
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits
; APPLICANT: Hoeben, Robert
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: Van der Eb, Alex J.
; APPLICANT: Schouten, Govert
; TITLE OF INVENTION: PACKAGING SYSTEMS
; FILE REFERENCE: 2578-393505
; CURRENT APPLICATION NUMBER: US/09/912,552
; CURRENT FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: US/09/356,575
; PRIOR FILING DATE: 1999-07-19
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 20
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; OTHER INFORMATION: Derived from Adenovirus
US-09-912-552-20
Query Match      100.0%; Score 55; DB 9; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.5e-12;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  GTACATGATGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACACTAGTCAAT 55
Db      1  GTACATGATGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACACTAGTCAAT 55
RESULT 3
US-09-918-029-20
; Sequence 20, Application US/09918029
; Patent No. US20020102732A1
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoeben, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: Van der Eb, Alex J.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO
; TITLE OF INVENTION: BE USED
; FILE REFERENCE: 3833.205
; CURRENT APPLICATION NUMBER: US/09/918,029
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: 09/506,548
; PRIOR FILING DATE: 2000-02-16
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 20
; LENGTH: 55
```

```
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: primer HP/c1a2
US-09-918-029-20
Query Match      100.0%; Score 55; DB 10; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.5e-12;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  GTACATGATGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACACTAGTCAAT 55
Db      1  GTACATGATGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACACTAGTCAAT 55
RESULT 4
US-10-038-271-20
; Sequence 20, Application US/10038271
; Patent No. US20020151032A1
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoeben, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: Van der Eb, Alex J.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO BE USED
; TITLE OF INVENTION: GENE THERAPY
; FILE REFERENCE: 3833.105
; CURRENT APPLICATION NUMBER: US/10/038,271
; CURRENT FILING DATE: 2001-10-23
; PRIOR APPLICATION NUMBER: 09/333,820
; PRIOR FILING DATE: 1999-06-15
; PRIOR APPLICATION NUMBER: US 08/793,170
; PRIOR FILING DATE: 1997-03-25
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 20
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: primer HP/c1a2
US-10-038-271-20
Query Match      100.0%; Score 55; DB 12; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.5e-12;
Matches 55; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  GTACATGATGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACACTAGTCAAT 55
Db      1  GTACATGATGACCTAGTGGCCGCCGGGCTTTGGCCGGGCGGCACACTAGTCAAT 55
RESULT 5
US-10-125-751-19/c
; Sequence 19, Application US/10125751
; Patent No. US20020173039A1
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoeben, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: Van der Eb, Alex J.
```

5 ATCGATTGACCTAGTGGCCCCGGGGCTTGGCCCCGGGGCAGTCTCAAT 55

```

; NAME/KEY: primer_bind
;
; LOCATION: (1)..(55)
;
; OTHER INFORMATION: /No. US20020119942A1e="primer HP/c1a1"
;

```

OTHER INFORMATION: Description of Artificial Sequence: primer
US-09-900-062-45

Query Match 92.7%; Score 51; DB 10; Length 55;
Best Local Similarity 100.0%; Pred. No. 5e-11;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 ATCGATTGACCTAGTGGCCGGGGCTTTGCCGGCGGCGACTAGGTCAAT 55
|||||
DB 55 ATCGATTGACCTAGTGGCCGGGGCTTTGCCGGCGGCGACTAGGTCAAT 5

RESULT 9
US-10-038-271-19/c
Sequence 19, Application US/10038271
Patent No. US20020151032A1

GENERAL INFORMATION:
APPLICANT: Fallaux, Fris J.
APPLICANT: Hoeber, Robert C.
APPLICANT: Bout, Abraham
APPLICANT: Valerio, Domenico

APPLICANT: Van der Eb, Alex J.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO BE USED IN
FILE REFERENCE: 3833.1US
CURRENT APPLICATION NUMBER: US/10/038,271
CURRENT FILING DATE: 2001-10-23
PRIOR APPLICATION NUMBER: 09/333,820
PRIOR FILING DATE: 1999-06-15
PRIOR APPLICATION NUMBER: US 08/793,170
PRIOR FILING DATE: 1997-03-25
PRIOR APPLICATION NUMBER: PCT/NL96/00244
PRIOR FILING DATE: 1996-06-14
PRIOR APPLICATION NUMBER: EP 95201728.3
PRIOR FILING DATE: 1995-06-26
PRIOR APPLICATION NUMBER: EP 95201611.1
PRIOR FILING DATE: 1995-06-15
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Corel WordPerfect 8.0
SEQ ID NO 19
LENGTH: 55
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
NAME/KEY:
LOCATION:
OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/c1a1

US-10-038-271-19

Query Match 92.7%; Score 51; DB 12; Length 55;
Best Local Similarity 100.0%; Pred. No. 5e-11;
Matches 51; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 ATCGATTGACCTAGTGGCCGGGGCTTTGCCGGCGGCGACTAGGTCAAT 55
|||||
DB 55 ATCGATTGACCTAGTGGCCGGGGCTTTGCCGGCGGCGACTAGGTCAAT 5

RESULT 10
US-10-125-751-17/c
Sequence 17, Application US/10125751
Patent No. US20020173039A1

GENERAL INFORMATION:
APPLICANT: Fallaux, Fris J.
APPLICANT: Hoeber, Robert C.
APPLICANT: Bout, Abraham
APPLICANT: Valerio, Domenico

APPLICANT: Van der Eb, Alex J.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO
TITLE OF INVENTION: BE USED
FILE REFERENCE: 3833.2US
CURRENT APPLICATION NUMBER: US/10/125,751

US-10-125-751-17

CURRENT FILING DATE: 2002-04-18
PRIOR APPLICATION NUMBER: 09/506,548
PRIOR FILING DATE: 2000-02-16
PRIOR APPLICATION NUMBER: PCT/NL96/00244
PRIOR FILING DATE: 1996-06-14
PRIOR APPLICATION NUMBER: EP 95201728.3
PRIOR FILING DATE: 1995-06-26
PRIOR APPLICATION NUMBER: EP 95201611.1
PRIOR FILING DATE: 1995-06-15
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Corel WordPerfect 8.0
SEQ ID NO 17
LENGTH: 50
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
NAME/KEY:
LOCATION:
OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/aspl
US-10-125-751-17

Query Match 78.9%; Score 43.4; DB 9; Length 50;
Best Local Similarity 97.8%; Pred. No. 4e-08;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 11 TGACCTAGTGGCCGGGGCTTTGCCGGCGGCGACTAGGTCAAT 55
|||||
DB 49 TGACCTAGTGGCCGGGGCTTTGCCGGCGGCGACTAGGTCAAT 5

RESULT 11
US-10-125-751-18
Sequence 18, Application US/10125751
Patent No. US20020173039A1

GENERAL INFORMATION:
APPLICANT: Fallaux, Fris J.
APPLICANT: Hoeber, Robert C.
APPLICANT: Bout, Abraham
APPLICANT: Valerio, Domenico
APPLICANT: Van der Eb, Alex J.
TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO
TITLE OF INVENTION: BE USED
FILE REFERENCE: 3833.2US
CURRENT APPLICATION NUMBER: US/10/125,751
CURRENT FILING DATE: 2002-04-18
PRIOR APPLICATION NUMBER: 09/506,548
PRIOR FILING DATE: 2000-02-16
PRIOR APPLICATION NUMBER: PCT/NL96/00244
PRIOR FILING DATE: 1996-06-14
PRIOR APPLICATION NUMBER: EP 95201728.3
PRIOR FILING DATE: 1995-06-26
PRIOR APPLICATION NUMBER: EP 95201611.1
PRIOR FILING DATE: 1995-06-15
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Corel WordPerfect 8.0
SEQ ID NO 18
LENGTH: 50
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
NAME/KEY:
LOCATION:
OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/aspl
US-10-125-751-18

Query Match 78.9%; Score 43.4; DB 9; Length 50;
Best Local Similarity 97.8%; Pred. No. 4e-08;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 11 TGACCTAGTGGCCGGGGCTTTGCCGGCGGCGACTAGGTCAAT 55
|||||
DB 6 TGACCTAGTGGCCGGGGCTTTGCCGGCGGCGACTAGGTCAAT 50

US-10-125-751-18

```
RESULT 12
US-09-912-552-17/c
; Sequence 17, Application US/09912552
; Publication No. US20020187553A1
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits
; APPLICANT: Hoeben, Robert
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: van der Eb, Alex
; APPLICANT: Schouten, Goyert
; TITLE OF INVENTION: PACKAGING SYSTEMS
; FILE REFERENCE: 2578-393505
; CURRENT APPLICATION NUMBER: US/09/912,552
; CURRENT FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: US/09/356,575
; PRIOR FILING DATE: 1999-07-19
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 17
; LENGTH: 50
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; OTHER INFORMATION: Derived from Adenovirus
US-09-912-552-17

Query Match          78.9%; Score 43.4; DB 9; Length 50;
Best Local Similarity 97.8%; Pred. No. 4e-08;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 11 TGACCTAGTGGCCGCCGGGCTTTGCGCGGCGGCACTAGTCAAT 55
      |||||||
Db 49 TGACCTAGTGGCCGCCGGGCTTTGCGCGGCGGCACTAGTCAAT 5

RESULT 13
US-09-912-552-18
; Sequence 18, Application US/09912552
; Publication No. US20020187553A1
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits
; APPLICANT: Hoeben, Robert
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: van der Eb, Alex
; APPLICANT: Schouten, Goyert
; TITLE OF INVENTION: PACKAGING SYSTEMS
; FILE REFERENCE: 2578-393505
; CURRENT APPLICATION NUMBER: US/09/912,552
; CURRENT FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: US/09/356,575
; PRIOR FILING DATE: 1999-07-19
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 18
; LENGTH: 50
; TYPE: DNA
; ORGANISM: Unknown
```

```
; FEATURE:
; OTHER INFORMATION: Derived from Adenovirus
US-09-912-552-18

Query Match          78.9%; Score 43.4; DB 9; Length 50;
Best Local Similarity 97.8%; Pred. No. 4e-08;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 11 TGACCTAGTGGCCGCCGGGCTTTGCGCGGCGGCACTAGTCAAT 55
      |||||||
Db 6 TGACCTAGTGGCCGCCGGGCTTTGCGCGGCGGCACTAGTCAAT 50

RESULT 14
US-09-918-029-17/c
; Sequence 17, Application US/09918029
; Patent No. US20020102732A1
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoeben, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: van der Eb, Alex J.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO
; TITLE OF INVENTION: BE USED
; TITLE OF INVENTION: IN GENE THERAPY
; FILE REFERENCE: 3833.2US
; CURRENT APPLICATION NUMBER: US/09/918,029
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: 09/506,548
; PRIOR FILING DATE: 2000-02-16
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 17
; LENGTH: 50
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/aspl
US-09-918-029-17

Query Match          78.9%; Score 43.4; DB 10; Length 50;
Best Local Similarity 97.8%; Pred. No. 4e-08;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 11 TGACCTAGTGGCCGCCGGGCTTTGCGCGGCGGCACTAGTCAAT 55
      |||||||
Db 49 TGACCTAGTGGCCGCCGGGCTTTGCGCGGCGGCACTAGTCAAT 5

RESULT 15
US-09-918-029-18
; Sequence 18, Application US/09918029
; Patent No. US20020102732A1
; GENERAL INFORMATION:
; APPLICANT: Fallaux, Frits J.
; APPLICANT: Hoeben, Robert C.
; APPLICANT: Bout, Abraham
; APPLICANT: Valerio, Domenico
; APPLICANT: van der Eb, Alex J.
; TITLE OF INVENTION: PACKAGING SYSTEMS FOR HUMAN RECOMBINANT ADENOVIRUS TO
; TITLE OF INVENTION: BE USED
; TITLE OF INVENTION: IN GENE THERAPY
; FILE REFERENCE: 3833.2US
; CURRENT APPLICATION NUMBER: US/09/918,029
```

```

; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: 09/506,548
; PRIOR FILING DATE: 2000-02-16
; PRIOR APPLICATION NUMBER: PCT/NL96/00244
; PRIOR FILING DATE: 1996-06-14
; PRIOR APPLICATION NUMBER: EP 95201728.3
; PRIOR FILING DATE: 1995-06-26
; PRIOR APPLICATION NUMBER: EP 95201611.1
; PRIOR FILING DATE: 1995-06-15
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Corel WordPerfect 8.0
; SEQ ID NO 18
; LENGTH: 50
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY:
; LOCATION:
; OTHER INFORMATION: Description of Artificial Sequence: PCT primer HP/asp2
US-09-918-029-18

```

```

Query Match          78.9%; Score 43.4; DB 10; Length 50;
Best Local Similarity 97.8%; Pred. No. 4e-08; 1; Indels 0; Gaps 0;
Matches 44; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

```

```

OY 11 TGACCTAGTGGCGCCGCGCTTTGCGCGGCGGCGCACTAGTCAAT 55
      |||||||
Db 6 TGACCTAGTGGCGCCGCGCTTTGCGCGGCGGCGCACTAGTCAAT 50

```

Search completed: December 27, 2002, 06:18:34
 Job time : 36 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 27, 2002, 04:38:01 ; Search time 1321.5 Seconds
(without alignments)
674,046 Million cell updates/sec

Title: US-09-918-029-20

Sequence: 1 gtccatcagttgacctagtg.....ccgggcgcgcactagtgcaat 55

Scoring table: IDENTITY_NUC
Gap 10.0 , Gape 1.0

Searched: 16154066 seqs, 809774376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :
1: em_estda:*
2: em_esthm:*
3: em_estin:*
4: em_estnu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_hlc:*
9: gb_estl:*
10: gb_estc2:*
11: gb_hlc:*
12: gb_estl3:*
13: gb_estl4:*
14: gb_estl5:*
15: em_estfun:*
16: em_estom:*
17: gd_gss:*
18: em_gss_hum:*
19: em_gss_inv:*
20: em_gss_pln:*
21: em_gss_vrt:*
22: em_gss_fun:*
23: em_gss_mam:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rod:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	24.8	45.1	756	10	BE254363 601109113
2	24.2	44.0	17	AZ205863	AZ205863 SP_0105_A
3	24.2	44.0	794	10	BE294341 601172854
4	23.8	43.3	174	17	A0588337 C1TB1-E1-
5	23.8	43.3	885	9	AL564683 AL564683
6	23.8	43.3	885	9	AL564683 AL564683

C	7	23.8	43.3	1108	14	B0887187	B0887187 AGENCOURT
C	8	23.6	42.9	319	17	A0357767	A0357767 C1TB1-E1-
C	9	23.6	42.9	372	13	B1716319	B1716319 1031009F0
C	10	23.2	42.2	409	13	B0955901	B0955901 CM3-CT060
C	11	23.2	42.2	977	14	B0672537	B0672537 AGENCOURT
C	12	23	41.8	317	14	R77680	R77680 y162c09..s1
C	13	23	41.8	596	12	BC194272	BC194272 RST13417
C	14	23	41.8	822	17	CNS021IX	CNS021IX Tetradon
C	15	23	41.8	906	17	CNS03A57	CNS03A57 Tetradon
C	16	23	41.8	1488	12	BC845293	BC845293 Tetradon
C	17	22.8	41.5	344	9	AA081549	AA081549 zn21907.r
C	18	22.8	41.5	453	17	TA313C10Q	TA313C10Q T. brucei
C	19	22.8	41.5	526	14	B0812739	B0812739 1030032A0
C	20	22.8	41.5	565	12	BE725257	BE725257 B64082806
C	21	22.8	41.5	592	9	AL703582	AL703582 DKF22686C
C	22	22.8	41.5	658	12	BC724427	BC724427 602693726
C	23	22.8	41.5	749	12	BC722376	BC722376 602693574
C	24	22.8	41.5	749	12	BC844791	BC844791 1024007G1
C	25	22.8	41.5	755	12	BC719715	BC719715 602690436
C	26	22.8	41.5	810	12	BC671517	BC671517 DRNBTE09
C	27	22.8	41.5	814	13	BC717933	BC717933 602693988
C	28	22.8	41.5	816	13	B1644399	B1644399 603204058
C	29	22.8	41.5	820	13	BC923596	BC923596 602823455
C	30	22.8	41.5	959	14	B0679905	B0679905 AGENCOURT
C	31	22.6	41.1	534	17	A0838837	A0838837 HS_4716_A
C	32	22.6	41.1	556	13	BM105647	BM105647 509147 MA
C	33	22.6	41.1	582	9	A1981396	A1981396 pat PK005
C	34	22.6	41.1	749	12	BC844791	BC844791 1024007G1
C	35	22.6	41.1	798	12	BE965525	BE965525 602125006
C	36	22.6	41.1	1051	9	AL549845	AL549845 AL549845
C	37	22.6	41.1	1115	13	BM546853	BM546853 AGENCOURT
C	38	22.6	41.1	1374	14	BM906040	BM906040 AGENCOURT
C	39	22.4	40.7	447	17	BC792158	BC792158 UTSW_R2E8
C	40	22.4	40.7	452	17	A0878057	A0878057 HS_2160_A
C	41	22.4	40.7	560	12	BE930291	BE930291 KC5-NT018
C	42	22.4	40.7	652	12	BE204631	BE204631 601867053
C	43	22.4	40.7	741	12	BC828205	BC828205 602753589
C	44	22.4	40.7	781	13	B1115162	B1115162 602863085
C	45	22.4	40.7	807	13	B1822108	B1822108 603039994

ALIGNMENTS

RESULT 1
BE254363
LOCUS 756 bp mRNA linear EST 13-JUL-2000
DEFINITION 601109113F1 NIH_MGC_16 Homo sapiens CDNA clone IMAGE:3350090 5',
mRNA sequence.
BE254363
ACCESSION BE254363.1 GI:9124791
VERSION
KEYWORDS
SOURCE
ORGANISM
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 756)
NIH-MGC <http://mgc.ncl.nih.gov/>.
AUTHORS
TITLE
JOURNAL
COMMENT
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LIML)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LIML at: image.llnl.gov
Plate: LHCMI45 row: 1 column: 03
High quality sequence stop: 668.
Location/Qualifiers
1..756
/organism="Homo sapiens"

FEATURES
source

```

/db_xref="taxon:9606"
/clone_image:3350090"
/clone_lib="NIH_MGC_16"
/tissue_type="retinoblastoma"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: eye; Vector: pOTB7; Site: 1: XhoI; site: 2:
EcoRI; cDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(G). Library constructed by Ling Hong
in the Laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC Library."

BASE COUNT      115 a      197 c      275 g      169 t
ORIGIN

Query Match      45.1%; Score 24.8; DB 10; Length 756;
Best Local Similarity 72.7%; Pred. No. 2.8e+02;
Matches 32; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

OY      8 GATTGACCTAGTGGCCCGGCTTTGGCCGGCGGCACTAGT 51
      11 11 11 11 11 11 11 11 11 11 11 11 11 11 11
Db      185 GACTGGCTGTGACCTGGGCGGTGCCGGGCGGCGCTGGGT 228

RESULT 2
LOCUS      A2205863      590 bp      DNA      linear      GSS 31-AUG-2000
DEFINITION      SE_0105_A1.P05.T7A Strongylocentrotus purpuratus, purple sea urchin
                  clone Plate=105 Col=9 Row=W.K. DNA sequence.
ACCESSION      A2205863
VERSION
KEYWORDS
SOURCE
ORGANISM
                  Strongylocentrotus purpuratus.
                  Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;
                  Echinoidea; Enechinoidea; Echinacea; Echinoida;
                  Strongylocentrotidae; Strongylocentrotus.
                  1 (bases 1 to 590)
REFERENCE
                  Cameron R.A., Mahiras G., Rast J.P., Martinez P., Biondi T.R.,
                  Swartzell S., Wallace J.C., Poustka A.J., Livingston B.T., Wray
                  G.A., Eftensohn C.A., Lehrach H., Britten R.J., Davidson E.H. and
                  Hood L.
                  A sea urchin genome project: Sequence scan, virtual map, and
                  additional resources
                  Proc. Natl. Acad. Sci. U.S.A. 97 (17), 9514-9518 (2000)
JOURNAL
MEDLINE      20402566
CONTACT      Cameron, RA, Davidson, EH, Hood, L
                  Division of Biology 156-29
                  California Institute of Technology
                  Pasadena California 91125, USA
                  Tel: (626) 395-8421
                  Fax: (626) 793-3047
                  Email: acameron@caltech.edu
                  Plate: 105 row: K column: 9
                  Seq primer: T7
                  Class: BAC ends
                  High quality sequence stop: 590.
FEATURES
Source
1..590
location/Qualifiers
1..590
/organism="Strongylocentrotus purpuratus"
/db_xref="taxon:7668"
/clone_image="105 Col=9 Row=W.K"
/clone_lib="Strongylocentrotus purpuratus, purple sea
  urchin, sperm genomic BAC library"
/notes="Organ: sperm; Vector: BACs 3.6; BAC Clones in E-Coli
  DH10B"

BASE COUNT      163 a      157 c      110 g      159 t      1 others
ORIGIN

Query Match      44.0%; Score 24.2; DB 17; Length 590;
Best Local Similarity 66.0%; Pred. No. 4.2e+02;

```

```

Matches 35; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

OY      3 ACATCATTCAGCTAGTGGCCCGGCGTTCGCCGGGCGGCACTAGTCAAT 55
      11 11 11 11 11 11 11 11 11 11 11 11 11 11 11
Db      165 ACATCACTACACACTGCTGCACAGGAGTACACTGTCCGCACTAGTCAAT 217

RESULT 3
LOCUS      BE294341/c      794 bp      mRNA      linear      EST 20-JUN-2000
DEFINITION      601172854F1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:3528349 5',
                  mRNA sequence.
ACCESSION      BE294341
VERSION
KEYWORDS
SOURCE
ORGANISM
                  human.
                  Homo sapiens
                  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                  Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
                  1 (bases 1 to 794)
                  NIH-MGC http://mgi.nci.nih.gov/.
                  National Institutes of Health, Mammalian Gene Collection (MGC)
                  Unpublished (1999)
                  Contact: Robert Strausberg, Ph.D.
                  Email: cgapbs@mail.nih.gov
                  Tissue Procurement: ATCC
                  cDNA Library Preparation: Ling Hong/Rubin Laboratory
                  cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
                  DNA Sequencing by: Incyte Genomics, Inc.
                  Clone distribution: MGC clone distribution information can be
                  found through the I.M.A.G.E. Consortium/LNL at: image.lnl.gov
                  Plate: LNCM197 row: 1 column: 14
                  High quality sequence stop: 605.
FEATURES
Source
1..794
location/Qualifiers
1..794
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_image="3528349"
/clone_lib="NIH_MGC_17"
/tissue_type="rhabdomyosarcoma"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: muscle; Vector: pOTB7; Site: 1: EcoRI;
  site: 2: XhoI; cDNA made by oligo-dT priming.
  Directionally cloned into EcoRI/XhoI sites using the
  following 5' adaptor: GGCACGAG(G). Size-selected >500bp
  for average insert size 1.8kb. Library constructed by
  Ling Hong in the Laboratory of Gerald M. Rubin (University
  of California, Berkeley) using ZAP-cDNA synthesis kit
  (Stratagene) and Superscript II RT (Life Technologies)."

BASE COUNT      193 a      226 c      216 g      159 t
ORIGIN

Query Match      44.0%; Score 24.2; DB 10; Length 794;
Best Local Similarity 71.1%; Pred. No. 4.5e+02;
Matches 32; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

OY      6 TCGATTGACCTAGTGGCCCGGCGTTCGCCGGGCGGCACTAGG 50
      11 11 11 11 11 11 11 11 11 11 11 11 11 11 11
Db      684 TCCCTGGGCTTAGTGGCCCTCGGGAGTGTCCCGGGCCATTGGG 640

RESULT 4
LOCUS      A0588337      174 bp      DNA      linear      GSS 07-JUN-1999
DEFINITION      CITBI-EI.2644P17.TR CITBI-EI Homo sapiens genomic clone 2644P17,
                  DNA sequence.
ACCESSION      A0588337
VERSION
KEYWORDS
SOURCE
ORGANISM
                  human.
                  Homo sapiens
                  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                  Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

```


REFERENCE	1 (bases 1 to 174)	
AUTHORS	Zhao,S., Adams,M.D., Niernan,W., Malek,J., Shizuya,H., Simon,M. and Venter,J.C.	
TITLE	Use of BAC End Sequences from CalTech Libraries for Sequence-Ready Map Building	
JOURNAL	Unpublished (1997)	
COMMENT	other.GSSs: CIBI-EI-2644P17.TF Department of Eukaryotic Genomics The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850 Tel: 301 838 0200 Fax: 301 838 0208 Email: hboettig@ig.org Clones are available from Research Genetics (info@resgen.com). BAC end search page: http://www.tlgr.org/tldb/humgen/bac_end_search/bac_end_search.html . Seq primer: M13 Reverse Class: BAC ends.	
FEATURES	Location/Qualifiers	
Source	1..174	
	/organism="Homo sapiens"	
	/db_xref="taxon:9606"	
	/clone="2644P17"	
	/clone_lib="CIBI-EI"	
	/sex="male"	
	/cell_type="sperm"	
	/note="Vector: pBeloBAC11; Site_1: EcoRI; Site_2: EcoRI; Caltech Human BAC Library D"	
BASE COUNT	47 a 47 c 49 g 30 t 1 others	
ORIGIN		
Query Match	43.3%; Score 23.8; DB 17; Length 174;	
Best Local Similarity	72.1%; Pred.No. 4.5e+02;	
Matches 31; Conservative	0; Mismatches 12; Indels 0; Gaps 0;	
QY	10	TTGACCTAGTGGCCGCCGCTTTGCCCGGCGGACATAGTGC 52
Db	76	TTGCCTAGTACTACCTGCGCTTCCCGGAGAGCAGAAGTGC 118
RESULT 5		
LOCUS	AL564683 885 bp mRNA linear EST 16-FEB-2001	
DEFINITION	AL564683 LTI_NFL001.NBC4 Homo sapiens cDNA clone CS0DM007YK12 3 prime, mRNA sequence.	
ACCESSION	AL564683	
VERSION	AL564683.1 GI:12915335	
KEYWORDS	EST.	
SOURCE	human.	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.	
AUTHORS	1 (bases 1 to 885)	
TITLE	L1,W.B., Gruber,C., Jesse,J. and Polayes,D.	
JOURNAL	Full-length cDNA libraries and normalization	
COMMENT	Unpublished (2001) Contact: Genoscope Genoscope - Centre National de Sequences BP 191 91006 Evry cedex - France Email: seque@genoscope.cns.fr, Web : www.genoscope.cns.fr .	
FEATURES	Location/Qualifiers	
Source	1..885	
	/organism="Homo sapiens"	
	/db_xref="taxon:9606"	
	/clone="CS0DM007YK12"	
	/clone_lib="LTI_NFL001.NBC4"	
	/sex="male"	
	/tissue_type="neuroblastoma cells"	
	/lab_host="DH10B"	
	/note="Organ: brain; Vector: pCMVSPORT 6; 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-stranded cDNA was digested with Not I and	

	<p>cloned into the Not I and Eco RV sites of the pCMVSPORT 6 vector. Library was normalized. Contact : Feng Liang Life Technologies, a division of Invitrogen 9800 Medical Center Drive Rockville, Maryland 20850, USA Fax : (1) 301 610 8371 Email : fliang@lifetech.com URL : http://fulllength.invitrogen.com"</p>					
BASE COUNT	166 a	253 c	263 g	159 t	44 others	
ORIGIN						
Query Match	Best Local Similarity	43.3%	Score 23.8;	DB 9;	Length 885;	
	Matches 31; Conservative	66.0%;	Pred. No. 6.2e+02;	Mismatches 14;	Indels 0;	Gaps 0;
OY	8 GATTGACCTAGTGC					54
Dd	561 GGTTCACACAGTGCCGCGAGTGTGACTGCGCGCACACKSCCKCA					607
RESULT 6						
LOCUS	AL564683/c	885 bp	mRNA	linear	EST 16-FEB-2001	
DEFINITION	AL564683 LTI_NFL001.NBC4 Homo sapiens CDNA clone CSODM007YK12 3					
ACCESSION	prime mRNA sequence.					
VERSION	AL564683					
KEYWORDS	AL564683.1 GI:12915335					
SOURCE	EST.					
ORGANISM	human.					
	Homo sapiens					
REFERENCE	Eukaryota; Metazoa; Chordata; Craniota; Vertebrata; Euteleostomi;					
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.					
TITLE	1 (bases 1 to 885)					
JOURNAL	Ll,W.B., Gruber,C., Jessee,J. and Polayes,D.					
COMMENT	Full-length cDNA libraries and normalization Unpublished (2001) Contact: Genoscope Genoscope - Centre National de Sequencage BP 191 91006 Evry cedex - France Email: seque@genoscope.cns.fr , Web : www.genoscope.cns.fr .					
FEATURES	Location/Qualifiers					
source	1..885					
	/organism="Homo sapiens"					
	/db_xref="taxon:9606"					
	/clone="CSODM007YK12"					
	/clone_1lb="LTI_NFL001.NBC4"					
	/sex="male"					
	/library_type="neuroblastoma cells"					
	/lab_host="DH10B"					
	/note="Organ: Brain; Vector: pCMVSPORT 6; 1st strand cDNA was primed with a NotI-Oligo(dT) primer. Five prime end enriched, double-stranded cDNA was digested with Not I and cloned into the Not I and Eco RV sites of the pCMVSPORT 6 vector. Library was normalized. Library was constructed by Life Technologies. Contact : Feng Liang Life Technologies, a division of Invitrogen 9800 Medical Center Drive Rockville, Maryland 20850, USA Fax : (1) 301 610 8371 Email : fliang@lifetech.com URL : http://fulllength.invitrogen.com "					
BASE COUNT	166 a	253 c	263 g	159 t	44 others	
ORIGIN						
Query Match	Best Local Similarity	43.3%	Score 23.8;	DB 9;	Length 885;	
	Matches 31; Conservative	66.0%;	Pred. No. 6.2e+02;	Mismatches 14;	Indels 0;	Gaps 0;
OY	8 GATTGACCTAGTGC					54
Dd	609 GGTTCACACAGTGCCGCGAGTGTGACTGCGCGCACACCTGTGCCAA					563
RESULT 7						
LOCUS	BQ887187/c	1108 bp	mRNA	linear	EST 16-AUG-2002	
	BQ887187					

DEFINITION AGENCOURT_8678059 NIH_MGC_40 Homo sapiens cDNA clone IMAGE:6380936
5', mRNA sequence.
ACCESSION BO887187
VERSION BO887187.1 GI:22279201
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 1108)
NIH-MGC <http://mgs.nci.nih.gov/>.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
CONTACT: Robert Strausberg, Ph.D.
Email: cgaps@email.nih.gov
Tissue Procurement: DCTD/DTF
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
<http://image.lnl.gov>
Plate: LNCM2570 row: f column: 09
High quality sequence start: 112
High quality sequence stop: 276.
FEATURES
source
1..1108
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:6380936"
/clone_1lb="NIH_MGC_40"
/tissue_type="carcinoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: prostate; Vector: pORF7; Site_1: XhoI;
Site_2: EcoRI; cDNA made by oligo-dT priming.
Directionally cloned into EcoRI/XhoI sites using the
following 5' adaptor: GGCACGAG(G). Library constructed by
Ling Hong in the laboratory of Gerald M. Rubin (University
of California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC Library."

BASE COUNT 197 a 363 c 388 g 160 t
ORIGIN

Query Match 43.3%; Score 23.8; DB 14; Length 1108;
Best Local Similarity 66.7%; Pred. No. 6.5e+02;
Matches 34; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

QY 4 CATGATTGACCTGTCGCGCGGCTTGCGCGGCGGCACACTGTCAA 54
Db 711 CATGACCCCGCCGACGCGACCGCGGCTTGCGCGCGGCACACTGTCAA 661

RESULT 8
AQ357767 319 bp DNA linear GSS 24-JAN-1999
LOCUS CITBI-El-2535A15.TR CITBI-El Homo sapiens genomic clone 2535A15,
DEFINITION DNA sequence.
ACCESSION AQ357767
VERSION AQ357767.1 GI:4184940
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 319)
Zhao, S., Adams, M.D., Nierman, W., Malek, J., Shizuya, H., Simon, M. and
Venter, A.C.
TITLE Use of BAC End Sequences from Caltech Libraries for Sequence-Ready
Map Building
JOURNAL Unpublished (1997)
COMMENT Other-GSSs: CITBI-El-2535A15.TR
Contact: Shaying Zhao, William Nierman, Mark Adams

Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbeel@igrg.org
Clones are available from Research Genetics (Info@resgen.com). BAC
end search page:
http://www.tigr.org/cdb/hungen/bac_end_search/bac_end_search.html.
Seq primer: M13 Reverse
Class: BAC ends.
FEATURES
source
1..319
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="2535A15"
/clone_1lb="CITBI-El"
/sex="male"
/cell_type="sperm"
/note="Vector: pBelBAC11; Site_1: EcoRI; Site_2: EcoRI;
Caltech Human BAC Library D"

BASE COUNT 64 a 86 c 93 g 76 t
ORIGIN

Query Match 42.9%; Score 23.6; DB 17; Length 319;
Best Local Similarity 64.8%; Pred. No. 5.9e+02;
Matches 35; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

QY 1 GTTACATGATGACCTGTCGCGCGGCTTGCGCGGCGGCACACTGTCAA 54
Db 92 GGAAATGATGATGTCCTCTCCACAGAAATTCGAGGATGAGTAA 145

RESULT 9
B1716319 372 bp mRNA linear EST 19-SEP-2001
LOCUS B1716319
DEFINITION 1031009P01.x1 C. reinhardtii CC-1690, Stress II (normalized),
B1716319
DEFINITION Lambda Zap II Chlamydomonas reinhardtii cDNA, mRNA sequence.
ACCESSION B1716319
VERSION B1716319.1 GI:15692014
KEYWORDS EST.
SOURCE Chlamydomonas reinhardtii.
ORGANISM Chlamydomonas reinhardtii.
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadales; Chlamydomonas.
1 (bases 1 to 372)
Grossman, A., Chang, C.-W., Davies, J., Harris, E., Hauser, C., Lefebvre,
P., McDermott, J.P., Shrago, J., Sillflow, C. and Stern, D.
TITLE Analyses of the Chlamydomonas reinhardtii genome: A Model,
JOURNAL Unpublished (2001)
Vascular Plants. Project: 1031
CONTACT: Charles Hauser
DCMB Box 91000
Duke University
Durham, NC 27708-1000
Tel: 919 613 8159
Fax: 919 613 8177
Email: chauser@duke.edu.
FEATURES
source
1..372
Location/Qualifiers
/organism="Chlamydomonas reinhardtii"
/strain="CC-1690 wild type mt+ 21gr"
/db_xref="taxon:3055"
/clone_1lb="C. reinhardtii CC-1690, Stress II (normalized)
", Lambda Zap II"
/note="Vector: pBluescript II SK-; Site_1: EcoRI; Site_2:
XhoI; Stress condition II library, constructed by John
Davies and Jeffrey McDermott, combines cDNAs from CC-1690
cells grown to mid-log phase in TAP (NH4+ - containing)
and shifted to TAP - NO3- (24hrs); H2 production
conditions (0, 12hr, 24hr) see Melis et al., (2000) Plant
Phys. 122: 127-135; TAP + H2O2 (1, 12, 24 hr); TAP +

BASE COUNT
ORIGIN

76 a 116 c 97 g 83 t

Query Match	42.98;	Score 23.6;	DB 13;	Length 372;
Best Local Similarity	64.88;	Pred. No. 6.1e+02;		
Matches 35; Conservative	0;	Mismatches 19;	Indels 0;	Gaps 0;

Oy 2 TACATCGATTGACCTAGTGCCGCCGGGCTTGCCTCCCGGGCAGTAAGTCAAT 55
 ||||| ||||| ||||| |||||
Db 234 TAAATCTACTCGGGTAATGCCCCCTGGGGCTTTGTTCGGGCACCTCCCGGGCAT 287

RESULT 10					
BG955901/c					
LOCUS	BG955901	409 bp	mRNA	linear	EST 12-JUN-2001
DEFINITION	Cm3-C70607-130201-747-f01 C70607 Homo sapiens CDNA,				mRNA sequence.
ACCESSION	BG955901				
VERSION	BG955901.1	GI:14374072			
KEYWORDS	EST.				
SOURCE	human.				
ORGANISM	Homo sapiens				

REFERENCE
AUTHORS

1 (bases 1 to 409)
Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R., Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.E., Goldman, G.H., Carvalho, A.F., Matsushima, A., Jalela, G.S., Simpson, D.H., Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeleel, C.V., O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.

TITLE Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
MEDLINE 20202663

COMMENT Contact: Simpson A.J.G.

Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-27064922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
<http://www.ludwig.org.br/scripts/gethtml2.pl?tl=CM3&ct=CM3-CT0607>
130201-747-f01&ct=2001-02-13&tt4=1)
Seq primer: puc 18 forward
High quality sequence start: 18
High quality sequence stop: 176.

BASE COUNT	86 a	113 c	136 g	74 t
ORIGIN				

	Query Match	42.2%	Score 23.2	DB 13	Length 409
	Best Local Similarity	70.5%	Pred. No. 8	4e+02	
	Matches 31, Conservative	0	Mismatches 13	Indels 0	Gaps 0
0Y	4	CATCGATTTCACCTGATGTCGCGCGGCTTTGGCCGCGCGCCACT	47		
Db	256	CAGGTTTCACCAAGGCCCATCGCTCTTCCTCCCTCGGCCCACT	213		

OY 4 CATGATTTCACCTAGTGGCCCCCGGGCTTTGCCGGGGGCACT 47
 || | | | | | | | | | | | | | | | | | | |
Db 256 CAGGTTTCACCAGGGCCCATCGGTCTTCCCCTGGCGCACT 213

[illegible]

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
1	(bases 1 to 977)	NIH-MGC http://mgc.ncl.nih.gov/ .	National Institutes of Health, Mammalian Gene Collection (MGC)	Unpublished (1999)
	Contact: Robert Strausberg, Ph.D.			

CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LINTL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LINTL at:
<http://image.llnl.gov>
Plate: LICM2409 row: c column: 18
High quality sequence stop: 405.

```

FEATURES
SOURCE
    Location/Qualifiers
        1..977
            /organism="Homo sapiens"
            /db_xref="taxon:9606"
            /clone_image="6256457"
            /clone_11b="NH_MGC_102"
            /tissue_type="epidermoid carcinoma, cell line"
            /lab_host="DH10B (phage-resistant)"
            /note="Organ: salivary gland; Vector: pOT7; Site_1: XhoI; Site_2: EcoRI; cDNA made by oligo-dT priming directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GCCACGAC(C). Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NH_MGC library."
BASE COUNT
    194 a      332 c      279 g      171 t      1 others
ORIGIN

```

Query Match	42.2%	Score 23.2;	DB 14;	Length 977;
Best Local Similarity	70.5%	Pred. NO. 1e+03;		
Matches 31; Conservative	0;	Mismatches 13;	Indels 0;	Gaps 0;

QY 2 TACATGATTACCCAGTCCCGCCGGGCTTTGCCCGGGGGCA 45
 | | | | | | | | | | | | | | | | | |
 Db 810 TTCGGCGCTCCCAAGTCCCGCCGGGCGGTGCGCCGGGACA 853

RESULT 12	
R77680	
LOCUS	
DEFINITION	317 bp mRNA
ACCESSION	R77680.y156c09.s1 Soares placenta M2HP Homo sapiens cDNA clone
VERSION	IMAGE:43824.3 similar to gp:011863 AMILORIDE-SENSITIVE AMINE OXIDASE (HUMAN), mRNA sequence.
	R77680 GI:852790

KEYWORDS		EST.	
SOURCE		human.	
ORGANISM		Homo sapiens	
		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
		Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.	
REFERENCE		1 (bases 1 to 317)	
AUTHORS		Hillier,L., Clark,N., Dubque,T., Elliston,K., Hawkins,M., Holman,	
		,M., Hultman,M., Kucaba,T., Le.M., Lennon,G., Marra,M., Parsons,J.,	
		Rifkin.L., Rohlfing.T., Soares.M., Tan,F., Trevasaki,E., Waterston	
		,R., Williamson,A., Wohldmann,P. and Wilson,R.	
TITLE		The WashU-Merck EST Project	
JOURNAL		Unpublished (1995)	
COMMENT		Contact: Wilsen RK Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel.: 314 286 1800 Fax: 314 286 1810 Email: est@wustl.wustl.edu	
		Insert Size: 858	
		High quality sequence starts: 1	
		High quality sequence stops: 1	
		Source: IMAGE Consortium, LNLN.	
		This clone is available royalty-free through LNLN ; contact the	
		IMAGE Consortium (info@image.lnl.gov) for further information.	
		Trace considered overall poor quality	
		Insert Length: 858 Std Error: 0.00	
		Seq primer: Promega -2lm13	
		High quality sequence stop: 1.	
FEATURES		Location/Qualifiers	
source		1..317	
		/organism="Homo sapieus"	
		/db_xref="GDB:553028"	
		/db_xref="taxon:9606"	
		/clone="IMAGE:143824"	
		/clone.lib="Soares Placenta Nb2HP"	
		/sex="Female"	
		/dev_stage="placenta obtained at birth (full term)"	
		/lab_host="DH10B (ampicillin resistant)"	
		/note="Organ: placenta; Vector: pRTV3D (Pharmacla) with a	
		modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st	
		strand cDNA was primed with a Not I - oligo(dT) primer [5'	
		ACTGTGAAGATTTCGCCGCACAGAATTTTCTTTTTTTT 3'] ,	
		double-stranded cDNA was ligated to Eco RI adaptors	
		(pharmacia), digested with Not I and cloned into the Not I	
		and Eco RI sites of the modified pRTV3 vector. Library	
		went through one round of normalization. Library	
		constructed by Bento Soares and M.Fatima Bonadio.	
BASE COUNT		54 a 68 c 110 g 56 t 29 others	
ORIGIN			
Query Match		41.8%; Score 23; DB 14; Length 317;	
Best Local Similarity		64.0%; Pred.No. 9.3e+07;	
Matches		Conservative 32; Conservative 0; Mismatches 18; Indels 0; Gaps 0;	
yqy		1 GTCAATCGATTGACACTAGTGCCGGCAGCTTTTGGCCGGCGCACCTAGG 50	
		I I I I I I I I I I I I I I I I	
Dd		12 GTTAATGNCNTGCCCCGAGTAGAGCCC GGCGGNWNCGGGAGAACATGGG 61	
RESULT 13			
BGI94272/c			
LOCUS		BGI94272	
DEFINITION		BSL13417 Atherys RAGE library Homo sapiens CDNA, mRNA sequence.	
ACCESION		BGI94272	
VERSION		BGI94272.1 GI:13715859	
KEYWORDS		EST.	
SOURCE		human.	
ORGANISM		Homo sapiens	
		Eukaryota; Metazoa; Chordata; Craniata; Vertebra; Euteleostomi;	
		Mammalia; Buthera; Primates; Catarrhini; Homlnidae; Homo.	
REFERENCE		1 (bases 1 to 596)	
AUTHORS		Harrington,J.J., Sherf,B., Rundlett,S., Jackson,P.D., Perry.R.,	
		Cain.S., Ievenhal,C., Thornton.M., Ramachandran,R., Whittington,J	

TITLE	JOURNAL	MEDLINE	COMMENT
'Lerner,L., Costanzo,D., McElligott,K., Booser,S., Mays,R., Smith , E., Veloso,N., Kikka,A., Hess,J., Cochran,K., Lo,K., Offenbacher ,J., Danzig,V. and Ducar,M. Creation of genome-wide protein expression libraries using random activation of genome-wide protein expression Nat. Biotechnol. 19 (5), 440-445 (2001) 21227151			Contact: Scott J. Cain Athersys, Inc. 3201 Carnegie Ave, Cleveland, OH 44115, USA Tel: 216 431 9900 Fax: 216 361 9596 Email: scaine@atersys.com High quality sequence stop: 596. Location/Qualifiers 1..596 /organism="Homo sapiens" /db_xref="taxon:9606" /clone_lib="Athersys RAGE Library" /cell_line="HT1080" /note="See 'Creation of Genome-wide Protein Expression libraries using Random Activation of Gene Expression', Nature Biotechnology, in press. Note that even though the cell type indicated is HT1080, since a random activation method was used, these sequence tags are not necessarily expressed in HT1080 under normal circumstances.'
BASE COUNT	181 a 140 c 126 g 148 t		
ORIGIN			
Query Match	41.8% Score 23; DB 12; Length 596;		
Best Local Similarity	68.1% Pred. No. 1.1e+03;		
Matches	32; Conservative 0; Mismatches 15; Indels 0; Gaps 0;		
Db	8 GATTGACCTAGTCCCGCGGGCTTTGCCGCGGCACTAGGTCAA 54 1 111 11111 1 11 111 111 11 11111 54 GTTGGCTTAGTGGCTCCTTCTCTGCGCAAGGCTAGACCTGGTCAA 8		
RESULT 14			
CNS0211X			
LOCUS			
DEFINITION	CNS0211X 822 bp DNA linear GSS 12-MAY-2000 Tetradon nigroviridis genome survey sequence PUC-Orl end of clone 225117 of library G from Tetradon nigroviridis, genomic survey sequence.		
ACCESSION	AL176946		
VERSION	AL176946.1 GI:7815003		
KEYWORDS	GSS: genome survey sequence.		
SOURCE	Tetradon nigroviridis.		
ORGANISM	Tetradon nigroviridis Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Acanthomorpha; Acanthopterygii; Percomorpha; Tetradontiformes; Tetradontidae; Tetradon.		
REFERENCE	1 (bases 1 to 822) Roest-Crolius,H., Jallou,O., Dasilva,C., Bouneau,L., Fisher,C., Bernot,A., Fizames,C., Wincker,P., Brottier,P., Queller,F., Saurin,W. and Weissenbach,J. Human gene number estimate provided by genome wide analysis using Tetradon nigroviridis DNA sequence		
TITLE	Unpublished		
JOURNAL	2 (bases 1 to 822)		
REFERENCE	Roest-Crolius,H., Jallou,O., Dasilva,C., Fizames,C., Fisher,C., Bouneau,L., Billault,A., Queller,F., Saurin,W., Bernot,A. and Weissenbach,J. Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetradon nigroviridis		
AUTHORS	Unpublished		
TITLE	3 (bases 1 to 822)		
JOURNAL	Genoscope.		
REFERENCE	Direct Submission		
AUTHORS	Submitted (12-Apr-2000)		
TITLE	This sequence is a single read and was generated as part of a large scale clone end sequencing project of the Tetradon nigroviridis		
COMMENT			

